

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets

(11)

EP 1 022 318 A2



(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
26.07.2000 Bulletin 2000/30

(51) Int. Cl.⁷: C09D 183/04, C08J 7/04,
C08K 3/00

(21) Application number: 00100911.7

(22) Date of filing: 18.01.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 19.01.1999 JP 1112199
21.05.1999 JP 14240999

(71) Applicant: JSR Corporation
Tokyo 104-8410 (JP)

(72) Inventors:
• Kanamori, Tarou,
c/o JSR Corporation
Tokyo 104-8410 (JP)

• Shimada, Mibuko,
c/o JSR Corporation
Tokyo 104-8410 (JP)
• Sakagami, Toshinori,
c/o JSR Corporation
Tokyo 104-8410 (JP)

(74) Representative:
Pellmann, Hans-Bernd, Dipl.-Ing. et al
Patentanwaltsbüro
Tiedtke-Bühling-Kinne & Partner
Bavariaring 4-6
80336 München (DE)

(54) **Method of making coating layers containing photocatalyst and a photocatalyst coating film formed thereby**

(57) A film which is improved in the resistance to fouling, the durability, and the transparency and has a photocatalytic function is provided by a method of making photocatalyst contained coating layers which comprises the steps of forming on the surface of an organic film an (under)coating of a coating composition composed mainly of at least one component (a) selected from organosilanes, hydrolyzates of the organosilanes, and condensates of the organosilanes and, when desired, a polymer component (b) having a silyl group containing silicon atoms bonded with a hydrolytic group and/or a hydroxy group, and forming on the (under)coating an (over)coating of a coating composition composed mainly of the component (a), a photocatalyst and, if desired, the component (b).

EP 1 022 318 A2

Description**Field of the Invention**

5 [0001] The present invention relates to a method of making photocatalyst contained coating layers to provide a film which is improved in the resistance to fouling, the durability, and the transparency while having a photocatalytic function, and to a photocatalyst coating film formed by the method.

Background of the Invention

10 [0002] It has lately been desired to provide a coating composition for forming a coating which is improved in the resistance to climate, the resistance to fouling, and the resistance to chemicals while being high in the physical hardness. It is also anticipated that such compositions should be arranged favorable for common use. Some conventional compositions composed mainly of a silyl-based vinyl resin and an organosilane compound are disclosed in Japanese Patent Laid-open Publications (Hei)01-69673 and (Hei)01-69674 where fundamental materials for making coatings are limited. Also, similar inventions are disclosed in Japanese Patent Laid-open Publications (Hei)04-108172 and (Hei)04-117473 where fundamental materials are widely selected but their details for coating performances are not definitely described.

15 [0003] Organosilane coating materials have technically been developed for use as maintenance-free coating materials which are advantageous for the resistance to climate (and sunlight), fouling, and so forth. The demand for improving the performance of organosilane coating materials is overwhelmingly increased. Therefore, it is now desired to provide coating materials which are improved in the coating appearance, the adhesivity, the resistance to climate, heat, alkalis, organic chemicals, moisture, and (hot) water, the insulating property, the abrasion resistance, the resistance to fouling, and so forth and are thus capable of forming a coating of a higher hardness.

20 [0004] Particularly for improving the resistance to fouling, it is known to make the surface of a coating hydrophilic. For example, the addition of a hydrophilic material or a water-soluble material is proposed. However, such a hydrophilic material or a water-soluble material may gradually be deteriorated when exposed to light and washed away with water, hence failing to maintain the level of hydrophilic property of the coating at the surface throughout a duration of service time.

25 [0005] Also, a variety of coating components including photocatalytic materials have been provided. One of them is, for example, a component for making a photocatalytic titan oxide coating composed mainly of titan oxide, hydrolyzate of a hydrolytic silicon compound (alkyl silicate or silicon halogenide), and a solvent (water or alcohol) (See Japanese Patent Laid-open Publication (Hei)8-164334). Another is a component treated at the surface for providing anti-bacteria and anti-mycotic properties and comprising a silicon compound having at least two alkoxy groups, a titan or zirconium compound having at least two alkoxy groups, and a hydrophilic non-organic powder e.g. of titan oxide processed with alkoxy silane and/or polysiloxane having at least two alkoxy groups (See Japanese Patent Laid-open Publication (Hei)8-176527). Moreover, a method is known for making a non-organic coating by producing a coating from a liquid mixture of a powder having a photocatalytic function and containing a non-organic paint which is prepared from 20 to 200 parts by weight of tetra-alkoxysilane, 100 parts by weight of tri-alkoxysilane, and from 0 to 60 parts by weight of dialkoxy silane and of which the weight-average molecular weight based on polystyrene is higher than 900 and processing the coating with an acid or an alkali (See Japanese Patent Laid-open Publication (Hei)8-259891).

30 [0006] However, the compositions for making a coating and the liquid mixture are intended only to provide anti-bacterial and anti-mycotic properties, remove odors, and decompose toxic substances in the alkoxy silane and/or polysiloxane containing a photocatalyst and a guanidyl group. In addition to the above, the above mentioned arts fail to 35 specify totally the essential characteristics of the resultant organosilane coating including the hardness, the adhesivity, and the resistance to alkalis, organic chemicals, climate, and fouling.

40 [0007] Further disclosed are a composition adapted for use as a cog composition which satisfies the requirements for performance of an organosilane coating material and composed mainly of a partial condensate of organosilane, an aqueous dispersion of colloidal silica, and a silicon-modified acrylic resin (See Japanese Patent Laid-open Publication (Sho)60-135465), a composition composed mainly of a condensate of organosilane, a chelate compound of zirconium alkoxide, and a hydrolytic silyl-based vinyl resin (Japanese Patent Laid-open Publication (Sho)64-1769), and a compound composed mainly of a condensate of organosilane, a colloidal alumina, and a hydrolytic silyl-based vinyl resin (See U.S. Patent No. 4,904,721).

45 [0008] However, coatings made of the compositions disclosed in (Sho)60-135465 and U.S. Patent No. 4,904,721 have the disadvantage that their glittering property may be declined when they are exposed to ultraviolet light for a considerable length of time. Also, the composition disclosed in (Sho)64-1769 is low in the storage stability; it may easily be turned to gel within a short period of time when its solid density is increased.

50 [0009] We, the inventors, have invented a composition for coating which includes a hydrolyzate and/or a partial con-

densate of organosilane, a vinyl resin having a silyl group containing silicon atoms bonded with a hydroxy group and/or a hydrolyzate, a metallic chelate compound, β -diketone and/or β -ketoester (See Japanese Patent Laid-open Publication (Hei)5-345877). While our composition has a favorable balance over the coating characteristics required for any organosilane coating material, more improvement of the coating characteristics shall be required.

5 [0010] Moreover, a self-cleaning film having a surface layer provided on a film substrate and containing substantially transparent particles of a photocatalytic oxide is disclosed in Japanese Patent Laid-open Publication (Hei)9-227161. The film may however be deteriorated by the effect of photocatalytic actions as being doubtful in the long-term durability. Also, a photocatalytic sheet having a photocatalytic coating formed on its upper surface and provided at its lower surface with an adhesive coating for attaching to another device or material to offer a photocatalytic function is
10 disclosed in Japanese Patent Laid-open Publication (Hei)9-313887. Similarly, the sheet may however be deteriorated by the effect of photocatalytic actions as being doubtful in the long-term durability.

Summary of the Invention

15 [0011] The present invention has been invented in view of overcoming the foregoing technical drawbacks of the prior arts and its object is to provide a method of making photocatalyst contained coating layers to provide a film which is improved in the resistance to fouling, the durability, and the transparency and has a photocatalytic function while the coating is easily applied on an organic film substrate, and provide a photocatalyst coating film produced by the method.

20 [0012] The method of making photocatalyst contained coating layers, according to the present invention, comprises the steps of providing a coating layer of the composition (i) or (ii) on the surface of an organic film and providing a coating layer of the composition (I) or (II) on the coating layer, wherein:

the composition (i) is a coating composition containing at least one component (a) selected from organosilanes, hydrolyzates of the organosilanes, and condensates of the organosilanes represented by Formula 1



(wherein, R^1 is a monovalent organic group having 1 to 8 carbon atoms: when two exist, they are either identical to or different from each other; R^2 is an alkyl group having 1 to 5 carbon atoms or an acyl group having 1 to 6 carbon atoms: when two exist, they are either identical to or different from each other; and n is an integer ranging from 0 to 2);

the composition (ii) is a coating composition containing the component (a) and a polymer component (b) having a silyl group having silicon atoms bonded with a hydrolytic group and/or a hydroxy group;

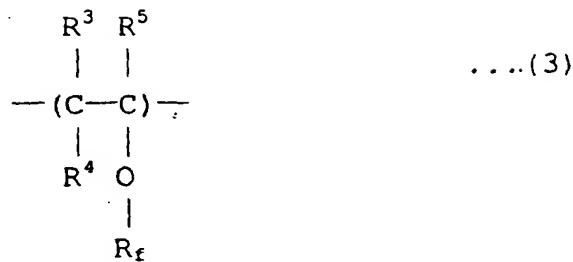
the composition (I) is a coating composition containing the component (a) and a photocatalyst (c); and

the composition (II) is a coating composition containing the component (a), the component (b), and the photocatalyst (c).

[0013] The component (b) in the composition (ii) may be at least a substance selected from polymers having a structural unit expressed by Formula 2



50 (wherein, R^3 to R^5 are C_mY_{2m+1} , m is an integer ranging from 0 to 5, and Y is selected from F, H, and Cl separately) and/or a structural unit expressed by Formula 3



(wherein, R_f is an alkyl group or an alkoxyalkyl group having fluorine atoms and R^3 to R^5 are analogous to those in Formula 2 or may be modified without departing from the term of analogy), and simultaneously having a silyl group having silicon atoms bonded with a hydrolytic group and/or a hydroxy group.

15

[0014] The component (b) in the composition (II) may be at least a substance selected from polymers having a structural unit expressed by Formula 2 and/or a structural unit expressed by Formula 3 and simultaneously having a silyl group containing silicon atoms bonded with a hydrolytic group and/or a hydroxy group.

20

[0015] The composition (i) or (ii) may further contain an ultraviolet light absorber (d).

[0016] The composition (i) or (II) may further contain a colloidal silica and/or a colloidal alumina (e).

[0017] The composition (i) or (ii) may further contain a colloidal silica and/or a colloidal alumina (e).

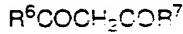
[0018] The component (a) in at least one of the compositions (i), (ii), (I), and (II) may partially or at its entirety have an SiO bond and a siloxane oligomer of which the weight average molecular weight ranges from 300 to 100,000.

25

[0019] Also, at least one of the compositions (i) (ii), (I), and (II) may preferably have a curing catalyst (f).

[0020] The curing catalyst (f) may preferably be an organic metal compound.

[0021] At least one of the compositions (i), (ii), (I), and (II) may preferably contain at least a compound selected from β -diketones and β -ketoesters (g) expressed by Formula 4



(4)

(wherein, R^5 is a monovalent hydrocarbon group having 1 to 6 carbon atoms, such as methyl, ethyl, n-propyl, i-propyl, n-butyl, sec-butyl, t-butyl, n-pentyl, n-hexyl, cyclohexyl, or phenyl, and R^7 is either a monovalent hydrocarbon group having 1 to 6 carbon atoms, like R^5 , or an alkoxy group having 1 to 16 carbon atoms, such as methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, sec-butoxy, t-butoxy, lauryloxy, or stearyoxy)

35

[0022] The photocatalyst coating film according to the present invention is produced by providing coating layers on the surface of an organic film with the use of a method of making photocatalyst contained coating layers defined above.

Detailed Description of the Preferred Embodiments

40

[0023] The present invention will now be described in a sequence.

Composition (I)

45

[0024] The composition (I) is a coating composition containing the following components (a) and (c).

Component (a)

50

[0025] The component (a) is at least a substance selected from organosilanes denoted by Formula 1 (referred to as organosilanes (1) hereinafter), hydrolyzates of the organosilanes (1), and condensates of the organosilanes (1), which acts as a bonding agent in the composition according to the present invention. More specifically, the component (a) may be of one of the three groups or a mixture of any two groups or a mixture of all the three groups.

55

[0026] Be noted that the hydrolyzate of an organosilane (1) is not limited to a particular one where all the OR^2 groups, generally two to four, in the organosilane (1) have been hydrolyzed but may be prepared where one or two or more of the groups have been hydrolyzed or may be a mixture of those groups.

[0027] The condensate of an organosilane (1) has an $Si-O-Si$ bond where the silanol group in a hydrolyzate of the organosilane (1) was condensed. It is not mandatory to condense all the silanol groups. The concept of a condensate of the organosilane (1) according to the present invention includes one where only a few of the silanol groups have been

condensed and a mixture of condensates whose levels of condensation are different.

[0028] In Formula 1, characteristic examples of the monovalent organic group of R^2 containing 1 to 8 carbon atoms are, for example, an alkyl group such as methyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, t-butyl, n-hexyl, n-heptyl, n-octyl, or 2-ethylhexyl; an acyl group such as acetyl, propionyl, butyryl, valeryl, benzoyl, trioyl, or caproyl; a vinyl group; an aryl group; a cyclohexyl group; a phenyl group; an epoxy group; a glycidyl group; a (meth-)acryloxy group; an ureide group; an amide group; a fluoroacetooamide group; and an isocyanate group as well as their substituent derivatives.

[0029] The substituent group in the substituent derivative of R^1 may be selected of a set of halogen atoms, a substituted or not-substituted amino group, a hydroxy group, a mercapto group, an isocyanate group, a glycidoxyl group, a 3,4-epoxycyclohexyl group, a (meth-)acryloxy group, an ureide group, and an ammonium base. The number of carbon atoms in the substituent derivative of R^1 including the carbon atoms in the substituent group is not greater than eight.

[0030] When two R^1 groups exist in Formula 1, they may be either identical or different.

[0031] The alkyl group of R^2 containing 1 to 5 carbon atoms may be, for example, methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, t-butyl, and n-pentyl. The acyl group may be, for example, acetyl, propionyl, butyryl, valeryl, and caproyl.

[0032] Two or more of the R^2 groups in Formula 1 may be either identical or different from each other.

[0033] Characteristic examples of the organosilane (1) are tetra-alkoxysilane including tetramethoxysilane, tetraethoxysilane, tetra-n-propoxysilane, tetra-i-propoxysilane, and tetra-n-butoxysilane; trialkoxysilane including methyl-trimethoxysilane, methyl triethoxysilane, ethyl-trimethoxysilane, n-propyl-trimethoxysilane, n-propyl-triethoxysilane, i-propyl-triethoxysilane, i-propyl-trimethoxysilane, i-propyl-triethoxysilane, n-butyl-trimethoxysilane, n-butyl-triethoxysilane, n-pentyl-trimethoxysilane, n-hexyl-trimethoxysilane, n-heptyl-trimethoxysilane, n-cetyltrimethoxysilane, vinyl-trimethoxysilane, vinyltriethoxysilane, cyclohexyl-trimethoxysilane, cyclohexyl-triethoxysilane, phenyl-trimethoxysilane, phenyl-triethoxysilane, 3-chloropropyl-trimethoxysilane, 3-chloropropyl-triethoxysilane, 3,3,3-trifluoropropyl-trimethoxysilane, 3,3,3-trifluoropropyl-triethoxysilane, 3-aminopropyl-trimethoxysilane, 3-aminopropyl-triethoxysilane, 2-hydroxyethyl-trimethoxysilane, 2-hydroxyethyl-triethoxysilane, 2-hydroxypropyl-trimethoxysilane, 2-hydroxypropyletriethoxysilane, 3-hydroxypropyl-trimethoxysilane, 3-hydroxypropyl-triethoxysilane, 3-mercaptopropyl-trimethoxysilane, 3-mercaptopropyl-triethoxysilane, 3-isocyanatepropyl-trimethoxysilane, 3-isocyanatepropyl-triethoxysilane, 3-glycidoxypropyl-trimethoxysilane, 3-glycidoxypropyl-triethoxysilane, 2-(3,4-epoxycyclohexyl) ethyl-trimethoxysilane, 2-(3,4-epoxycyclohexyl) ethyl-triethoxysilane, 3-(meth)acryloxypropyl-trimethoxysilane, 3-(meth)acryloxypropyl-triethoxysilane, 3-ureidepropyl-trimethoxysilane and 3-ureidepropyl-triethoxysilane; dialkoxy silanes including dimethyl-dimethoxysilane, dimethyl-diethoxysilane, diethyl-dimethoxysilane, di-n-propyl-dimethoxysilane, di-n-propyl-diethoxysilane, di-i-propyl-dimethoxysilane, di-i-propyl-diethoxysilane, di-n-butyl-dimethoxysilane, di-n-butyl-diethoxysilane, di-n-pentyl-dimethoxysilane, di-n-pentyl-diethoxysilane, di-n-hexyl-dimethoxysilane, di-n-hexyl-diethoxysilane, di-n-heptyl-dimethoxysilane, di-n-heptyl-diethoxysilane, di-n-octyl-dimethoxysilane, di-n-octyl-diethoxysilane, di-n-cyclohexyl-dimethoxysilane, di-n-cyclohexyl-diethoxysilane, diphenyl-dimethoxysilane, and diphenyl-diethoxysilane; methyl-triacetyloxy silanes; and dimethyl-diacetyloxy silanes.

[0034] Preferably, it may be selected from the trialkoxysilanes and the dialkoxy silanes. The trialkoxysilane may preferably be methyl-trimethoxysilane, methyl-triethoxysilane, 3-glycidoxypropyl-trimethoxysilane, 3-glycidoxypropyl-triethoxysilane, 3-(meth)acryloxypropyl-trimethoxysilane, or 3-(meta)acryloxypropyl-triethoxysilane. The dialkoxy silane may preferably be dimethyl-dimethoxysilane or dimethyl-diethoxysilane.

[0035] According to the present invention, the organosilane (1) may most preferably be trialkoxysilane or a combination of 40 to 95 mol% of trialkoxysilane and 60 to 5 mol % of dialkoxy silane. The use of dialkoxy silane with trialkoxysilane causes the resultant coating layer to be softened and increased in the resistance to alkalis.

[0036] The organosilane (1) may directly be used as a hydrolyzate and/or a condensate. The organosilane (1) employed as the hydrolyzate and/or the condensate enables to be utilized as the component (a) when it has been hydrolyzed and condensed. It is more preferable to perform the hydrolytic and condensing action of the organosilane (1) to yield the component (a) by feeding a proper amount of water during the preparation of the composition as a mixture of the organosilane (1) and the other components, as will explicitly be explained later.

[0037] According to the present invention, the component (a) may be provided as a single substance or a mixture of two or more substances.

[0038] The component (a) may either partially or totally include a siloxane oligomer which has SiO bonds and of which the weight-average molecular weight ranges from 300 to 100,000.

[0039] The siloxane oligomer may be provided as a single oligomer or a mixture of two or more oligomers. The ratio of O to Si in the SiO bond is normally from 0.5:1 to 2:1. The oligomer as a single substance may have a ratio of 1:1 (a 2-functional oligomer), 1.5:1 (a 3-functional oligomer), and 2:1 (a 4-functional oligomer). Most preferable is a siloxane oligomer in which the ratio of O to Si is 1.5:1 or 1:1. The method of producing the siloxane oligomer is of no limitations but may preferably be a process for condensation of the chlorosilane or alkoxy silane.

[0040] The end functional group of siloxane in the siloxane oligomer may be selected from a hydroxy group, a set of halogen atoms, an organic group having 1 to 15 carbon atoms (for example, alkyl, acyl, alkoxy, alkoxy silyl, vinyl, aryl,

acetoxy, acetoxyisilyl, cycloalkyl, phenyl, glycidyl, (meth)acryloxy, ureide, amide, fluoroacetoamide, or isocyanate), and a group expressed by $\text{Si}-(\text{RO})_p-(\text{R}'\text{O})_q-\text{R}''$ (wherein, R and R' are alkyl groups which are identical or difference, each having 1 to 5 carbon atoms, R'' is an alkyl radical having 1 to 5 carbon atoms or hydrogen atoms, and p and q are integers whereas p+q ranges from 2 to 30). Also, the groups may have been hydrolyzed and condensed partially or totally or may be substituent derivatives.

5 [0041] The set of halogen atoms may associate with fluorine and chlorine atoms.

[0042] The alkyl group having 1 to 15 carbon atoms may be selected from methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, sec-butyl, t-butyl, n-hexyl, n-heptyl, n-octyl, n-nonyl, n-decyl, undecyl, dodecyl, tridecyl, tetradeccyl, hexadecyl, heptadecyl, octadecyl, and 2-ethylhexyl. The acyl group may be selected from acetyl, propionyl, butyryl, valeryl, benzoyl, and trioyl. The alkoxy group may be selected from methoxy, ethoxy, propoxy, and butoxy. The alkoxyisilyl group may be selected from methoxyisilyl, ethoxyisilyl, propoxysilyl, and butoxysilyl. Also, vinyl, aryl, acetoxy, acetoxyisilyl, cyclohexyl, phenyl, glycidyl, (meth)acryloxy, ureide, amide, fluoroacetoamide, isocyanate, and their substituent derivatives may equally be used.

[0043] The group of $\text{Si}-(\text{RO})_p-(\text{R}'\text{O})_q-\text{R}''$ is a polyoxyalkylene group such as polyoxyethylene, polyoxypropylene, and poly(oxyethylene/oxypropylene). Because the siloxane oligomer has such end functional groups, a co-condensate of the component (a) and the siloxane oligomer is hydrophilic and simultaneously, the polyoxyalkylene groups is very likely to be absorbed by the other components (c) and (d) which in turn improve the dispersion stability.

[0044] The substituent group in the substituent derivative may be a set of halogen atoms, a substituted or not-substituted amino group, a hydroxy group, a mercapto group, an isocyanate group, glycidoxyl group, a 3,4-epoxycyclohexyl group, a (meth)acryloxy group, an ureide group, an ammonium base, or a ketoester group.

[0045] The weight-average molecular weight of the siloxane oligomer ranges from 300 to 100,000, preferably from 600 to 100,000, more preferably from 800 to 100,000, and most preferably from 1,000 to 50,000. If the level is smaller than 300, the resiliency of the resultant coating layer will decrease. If greater than 100,000, the composition in the coating layer will unfortunately be declined in the storage stability.

[0046] According to the present invention, the siloxane oligomer may be a mixture of two oligomers whose Mw levels are different from each other. For example, one oligomer has Mw=400 to 2,800 while the other has Mw=3,000 to 50,000.

[0047] The amount of the component (a) based on the total amount of a structural unit denoted by $(\text{R}^1_{2-n}\text{SiO}_{(4-n)/2})$ according to the present invention is preferably from 5 to 95 percent by weight and more preferably from 10 to 90 percent by weight; the same of the siloxane oligomer is preferably from 5 to 95 percent by weight and more preferably from 10 to 90 percent by weight (provided that (a)+siloxane oligomer=100 percent by weight). When the component (a) is smaller than 5 percent by weight, the composition in the resultant coating layer will be poor in the curing property. When higher than 95 percent by weight, the resistance to cracking will be declined.

[0048] The siloxane oligomer in the composition of the present invention is normally hydrolyzed and condensed together with the component (a).

[0049] In that case, the requirement of water for hydrolysis and condensation of the organosilane (1) and the siloxane oligomer is generally 0.3 to 3 moles and preferably 0.4 to 2 moles, based on one mole of the alkoxy groups.

[0050] The hydrolysis and condensation or the reaction during the synthesis of the composition of the present invention is carried out at a temperature of 30 to 80 °C, preferably 40 to 70°C, and for a duration of 0.5 to 10 hours, preferably 1 to 7 hours.

[0051] In the composition of the present invention, the weight-average molecular weight of a co-hydrolyzed condensate (a hydrolyzate and/or a condensate) of the component (a) and the siloxane oligomer is generally 500 to 100,000 and preferably about 600 to about 80,000.

[0052] The component (a) (which may contain a siloxane oligomer) is commercially available as MKC silicate manufactured by Mitsubishi Chemical, an ethyl silicate manufactured by Colcoat, a silicon resin manufactured by Toray/Dow-Corning, a silicon resin manufactured by Toshiba Silicon, a silicon resin manufactured by Shin-etsu Chemical Industries, a hydroxyl contained polydimethylsiloxane manufactured by Dow-Corning Asia, and a silicon oligomer manufactured by Nippon Unica. In the present invention, those products may be used directly or after subjected to the condensation.

50

Component (c)

[0053] The component (c) is a photocatalyst which is preferably a powder and/or a sol form of a semiconductor having a photocatalytic function.

[0054] Characteristic examples of the semiconductor having a photocatalytic function are TiO_2 , TiO_3 , SrTiO_3 , FeTiO_3 , WO_3 , SnO_2 , Bi_2O_3 , In_2O_3 , ZnO , Fe_2O_3 , RuO_2 , CdO , CdS , CdSe , GaP , GaAs , CdFeO_3 , MoS_2 , LaRhO_3 , GaN , CdP , ZnS , ZnSe , ZnTe , Nb_2O_5 , ZrO_2 , InP , GaAsP , InGaAlP , AlGaAs , PbS , InAs , PbSe , and InSb . Preferably it is TiO_2 containing the anatase crystalline structure or ZnO and more preferably TiO_2 .

[0055] It was found according to the present invention that the photocatalytic function of the component (c) caused the surface of the coating layer to turn to hydrophilic within a short time after exposed to lower light, thus significantly improving the resistance to fouling of the coating layer while not substantially impairing the other essential characteristics of the coating layer. Also, the component (c) in the composition (I) of the coating according to the present invention is commonly bonded with the component (a), hence ensuring the hydrophilic property and the resistance to fouling in the coating layer throughout a long period of time.

[0056] The semiconductor is desired to be used in the form of powder and/or sol. More particularly, the form is preferably selected from three different types: a powder, an aqueous sol as dispersed in water, and a solvent sol as dispersed in a polar solvent such as isopropyl alcohol or a non-polar solvent such as toluene. The solvent sol may be further diluted with water or the solvent depending on the degree of dispersion of the semiconductor. The smaller the average diameter of particles of the semiconductor in the given state, the better the photocatalytic function is implemented. When the average particle diameter of the semiconductor exceeds 0.3 μm , the shading action of the semiconductor will readily make the coating layer opaque. When smaller than 0.3 μm , the coating layer will remain transparent. Accordingly, the average particle diameter of the semiconductor can desirably be determined corresponding to the application of the composition.

[0057] In case that the component (c) is in the form of an aqueous sol or a solvent sol, the solids concentration is preferably not higher than 50 percent by weight and more preferably not higher than 40 percent by weight.

[0058] For feeding the component (c) into the composition, it is desired to add the component (c) to a combination of the component (a), the solvent, and any of other components (d) to (h) as desired. Alternatively, after the component (c) is fed during the preparation of the composition, the component (a) is subjected to the hydrolysis and the condensation under the presence of the component (c). The feeding of the component (c) during the preparation of the composition allows a semiconductor constituent in the component (c) to be condensed together with the component (a), whereby the durability of the resultant coating layer will be improved throughout a long period of time. Also, in case that the component (c) is an aqueous sol, it is preferably fed during the preparation of the composition. If the viscosity is high due to the presence of a component (g), which will be explained later, the component (c) may preferably be fed during the preparation of the composition. Moreover, in case that the composition of the present invention is provided as an enamel including colorants, the adjustment for a color may be conducted after the supply of the component (c) or feeding the component (c) and the colorants may be carried out at once.

[0059] According to the present invention, the component (c) may be provided as a single substance or a combination of two or more substances.

[0060] The amount of the component (c) to be used is generally 1 to 500 parts by weight in its solid state, based on 100 parts by weight of the structural unit $(\text{R}^1)_n\text{SiO}_{(4-n)/2}$ in the component (a) (wherein, R^1 and n conform to those in Formula 1), and preferably 5 to 400 parts by weight. If the amount is smaller than 1 part by weight, the resistance to fouling derived from the photocatalytic action will be declined. When the amount exceeds 500 parts by weight, the forming of the coating layer will be discouraged.

[0061] The composition (I) of the present invention consists mainly of the two essential components (a) and (c) and, if desired, may be added with other appropriate components which will be explained later. The composition (I) is commonly provided with its components dissolved or dispersed in a solvent.

[0062] For preparing the composition, water is used to carry out the hydrolysis and condensation of the organosilane (1) or disperse the colloidal components which will be explained later. The requirement of water according to the present invention is generally 0.3 to 3 moles and preferably 0.4 to 2 moles, based on one mole of the alkoxy group in the component (a).

[0063] According to the present invention, an organic solvent is used to assist uniform mixture of the component (a), the component (c), and the other relevant components to have a desired concentration of all solids in the composition. In addition, the organic solvent may assist each of a variety of coating processes and thus increase the dispersion stability and the storage stability of a resultant coating.

[0064] The organic solvent for assisting to mix up the components uniformly is of no limitations and may be selected from, for example, alcohols, aromatic hydrocarbons, ethers, ketones, and esters.

[0065] The alcohols for the organic solvent include specifically methanol, ethanol, n-propyl alcohol, i-propyl alcohol, n-butyl alcohol, sec-butyl alcohol, t-butyl alcohol, n-hexyl alcohol, n-octyl alcohol, ethylene glycol, diethylene glycol, triethylene glycol, ethylene glycol monobutyl ether, ethylene glycol monoethyl ether acetate, diethylene glycol monoethyl ether, propylene glycol monomethyl ether, propylene monomethyl ether acetate, and diacetone alcohol.

[0066] The aromatic hydrocarbons include specifically benzene, toluene, and xylene. The ethers include specifically tetrahydrofuran and dioxane. The ketones include specifically acetone, methylethyl ketone, methylisobutyl ketone, and di-isobutyl ketone. The esters include specifically ethyl acetate, propyl acetate, butyl acetate, and propylene carbonate.

[0067] The organic solvent may be provided as a single solvent or a combination of two or more solvents.

Composition (II)

[0068] The composition (II) consists mainly of the two essential components (a) and (c) plus the following component (b).

5

Component (b)

[0069] The component (b) is a polymer (referred to as a silyl based polymer hereinafter) where a silyl group having silicon atoms bonded with a hydrolytic group and/or a hydroxy group (referred to as a specific silyl group hereinafter) is linked to the end and/or side of a molecular chain of the polymer. The component (b) in the composition (II) allows the hydrolytic radical and/or hydroxy radical in its specific silyl group to be condensed together with the component (a) thus contributing to the improvement of the coating layer characteristics.

[0070] The amount of silicon in the component (b) is generally 0.001 to 20 percent by weight and preferably 0.01 to 15 percent by weight, based on the total amount of the component (b).

[0071] A preferable form of the specific silyl group is expressed by Formula 5:

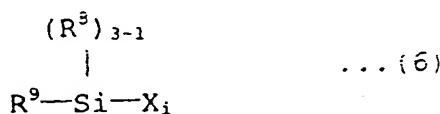


(wherein, X is a hydrolytic group or hydroxy groups of halogen atoms, an alkoxy group, an acetoxy group, a phenoxy group, a tioalkoxy group, or an amino group, R⁸ is hydrogen atoms, an alkyl group having 1 to 10 carbon atoms, or an aralkyl group having 1 to 10 carbon atoms, and i is an integer ranging from 1 to 3).

[0072] The component (b) may be produced by the following manners A and B.

[0073] Manner A: A method of adding a hydrosilane compound expressed by Formula 5 (referred to as a hydrosilane compound A hereinafter) to a carbon-carbon double bond in a carbon-carbon double bond containing vinyl polymer (referred to as an unsaturated vinyl polymer hereinafter).

[0074] Manner B: A method of co-polymerizing a silane compound (referred to as an unsaturated silane compound B hereinafter) which is expressed by Formula 6:



40 (wherein, X, R⁸, and i are identical to those in Formula 5 and R⁹ is an organic group having a polymeric double bond) with other vinyl monomers.

[0075] The hydrosilane compound A used in the manner A may be selected from, for example, halogenated silanes including methyl dichlorosilane, trichlorosilane, and phenyl dichlorosilane; alkoxy silanes including methyl dimethoxysilane, methyl diethoxysilane, phenyl dimethoxysilane, trimethoxysilane, and triethoxysilane; acyloxy silanes including methyl diacetoxysilane, phenyl diacetoxysilane, and triacetoxysilane; and aminoxy silanes including methyl dianinoxy silane, triaminoxy silane, and dimethyl aminoxy silane.

[0076] The hydrosilane compound A may be provided as a single compound or a combination of two or more compounds.

[0077] The unsaturated vinyl polymer having hydroxy groups and used in the manner A and having is of no limitations and may, for example, be produced by the following manners A1 and A2 and their combination.

[0078] Manner A1: A method of (co-)polymerizing vinyl monomers containing a functional group (referred to as a functional group α hereinafter) to yield a (co-)polymer and react the functional group α in the (co-)polymer with the carbon-carbon double bond contained unsaturated compound, which has a functional group (referred to as a functional group β hereinafter) reactable with the functional group α to yield an unsaturated vinyl polymer where the carbon-carbon double bond is linked to a side of the molecular chain of the polymer.

[0079] Manner A2: A method of (co-)polymerizing vinyl monomers with the use of either a radical polymerization initiator having the functional group α (for example, 4,4-azobis-4-cyanovalerianic acid) or a compound of a radical polymerization initiator and a chain transfer agent both having the functional group α (for example, 4,4-azobis-4-cyano-

valeric acid and dithioglycol acid) to yield a (co-)polymer where the functional groups α derived from the radical polymerization initiator and the chain transfer agent are linked to one or both ends of a molecular chain of the polymer, and causing the functional groups α in the (co-)polymer to react on the carbon-carbon double bond contained unsaturated compound having the functional group β to yield an unsaturated vinyl polymer where the carbon-carbon double bond is linked to one or both ends of a molecular chain of the polymer.

[0080] The reaction between the two functional groups α and β in the two manners A1 and A2 may involve esterification between a carboxyl group and a hydroxy group, ring opening esterification between a carbonic acid anhydrous group and a hydroxy group, ring opening esterification between a carboxyl group and an epoxy group, amidation between a carboxyl group and an amino group, ring opening amidation between a carbonic acid anhydrous group and an amino group, ring opening addition reaction between an epoxy group and an amino group, urethanation between a hydroxy group and an isocyanate group, or their combination.

[0081] The vinyl monomer having the functional group α may be selected from unsaturated carboxylic acid anhydrides including (meth)acrylic acid, crotonic acid, maleic acid, fumaric acid, and itaconic acid; hydroxy contained vinyl monomers including 2-hydroxyethyl(meth)acrylate, 2-hydroxypropyl(meth)acrylate, 3-hydroxypropyl(meth)acrylate, N-methyl(meth)acrylamide, and 2-hydroxyethylvinylether; amino contained vinyl monomers including 2-aminoethyl(meth)acrylate, 2-aminopropyl(meth)acrylate, 3-aminopropyl(meth)acrylate, and 2-aminoethylvinylether; amine-imide contained vinyl monomers including 1,1,1-trimethylamine8meth)acrylimide, 1,1-dmethyl-1-(2-hydroxypropyl)amine(meth)acrylimide, 1,1-dimethyl-1-(2'-phenyl-2'-hydroxyethyl)amine(meth)-acrylimide, and 1,1-dimethyl-1-(2'-hydroxy-2'-phenoxypropyl)amine(meth)acrylimide; and epoxy contained vinyl monomers including glycidyl(meth)acrylate and arylglycidylether.

[0082] The vinyl monomer having the functional group α may be provided in the form of a single monomer or a combination of two or more monomers.

[0083] The vinyl monomer capable of co-polymerizing with the functional group α containing vinyl monomer may be selected from:

- (A) aromatic vinyl monomers including styrene, α -methylstyrene, 4-methylstyrene, 2-methylstyrene, 3-methylstyrene, 4-methoxystyrene, 2-hydroxymethylstyrene, 4-ethylstyrene, 4-ethoxystyrene, 3,4-dimethylstyrene, 3,4-diethylstyrene, 2-chlorostyrene, 3-chlorostyrene, 4-chloro-3-methylstyrene, 4-t-butylstyrene, 2,4-dichlorostyrene, 2,6-dichlorostyrene, and 1-vinylphthalene;
- (B) (meth)acrylates including methyl(meth)acrylate, ethyl(meth)acrylate, propyl(meth)acrylate, α -butyl(meth)acrylate, β -butyl(meth)acrylate, amyl(meth)acrylate, β -amyl(meth)acrylate, hexyl(meth)acrylate, 2-ethylhexyl(meth)acrylate, α -octyl(meth)acrylate, and cyclohexyl-methacrylate;
- (C) multi-functional monomers including divinylbenzene, ethyleneglycol-di(meth)acrylate, diethyleneglycol-di(meth)acrylate, triethyleneglycol-di(meth)acrylate, tetraethyleneglycol-di(meth)acrylate, propyleneglycol-di(meth)acrylate, dipropyleneglycol-di(meth)acrylate, tripropyleneglycol-di(meth)acrylate, tetrapropyleneglycol-di(meth)acrylate, butanedioidi(meth)acrylate, hexanediol-di(meth)acrylate, trimethyloipropane-tri(meth)acrylate, and pentaerythritol-tetra(meth)acrylate;
- (D) acidic amide compounds including (meth)acrylamide, N-methylo(meth)acrylamide, N-methoxymethyl(meth)acrylamide, N-butoxymethyl(meth)acrylamide, N,N'-methylenebisacrylamide, diacetoneacrylamide, maleic acid amide, and maleimide;
- (E) vinyl compounds including vinyl chloride, vinylidene chloride, and fatty vinylester;
- (F) aliphatic conjugate dienes including 1,3-butadiene, 2-methyl-1,3-butadiene, 2,3-dimethyl-1,3-butadiene, 2-neopentyl-1,3-butadiene, 2-chloro-1,3-butadiene, 2-cyano-1,3-butadiene, isoprene, substituent straight-chain conjugate pentadiene substituted with a substituent group such as an alkyl group, halogen atoms, or a cyano group, and straight-chain or side-chain conjugate hexadiene;
- (G) vinyl cyanide compounds including acrylonitril and methacrylonitril;
- (H) fluorine atom containing monomers including trifluoroethyl(meth)acrylate and pentadecafluoroctyl(meth)acrylate;
- (I) piperidine monomers including 4-(meth)acryloiloxy-2,2,6,6-tetramethylpiperidine, 4-(meth)acryloiloamin-2,2,6,6-tetramethylpiperidine, and 4-(meth)acryloiloxy-1,2,2,6,6-pentamethylpiperidine; and other substances including caprolactam.

[0084] This type of the monomers may be provided in the form of a single monomer or a combination of two or more monomers.

[0085] The unsaturated compounds having the functional group α and the carbon-carbon double bond may be selected from vinyl monomers including a functional group α containing vinyl monomer, and isocyanate containing unsaturated compounds prepared by causing the hydroxy containing vinyl monomer to react on a di-isocyanate compound in equivalent mole relationship.

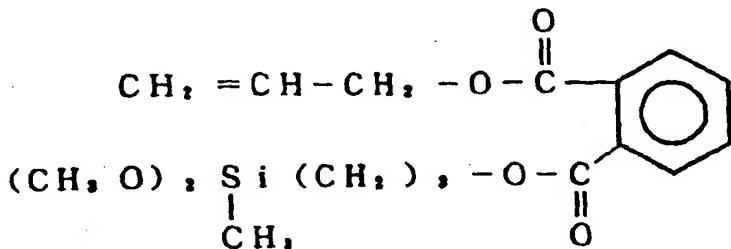
EP 1 022 318 A2

[0086] Characteristic examples of the unsaturated silane compound B used in the above described manner B are:

CH₂=CHSi(CH₃)(OCH₃)₂, CH₂=CHSi(OCH₃)₃, CH₃=CHSi(CH₃)Cl₂, CH₂=CHSiCl₃,
 CH₂=CHCOO(CH₂)₂Si(CH₃)₂(OCH₃)₂, CH₂=CHCOO(CH₂)₂Si(OCH₃)₃, CH₂=CHCOO(CH₂)₃Si(CH₃)(OCH₃)₂,
 CH₂=CHCOO(CH₂)₃Si(OCH₃)₃, CH₂=CHCOO(CH₂)₂Si(CH₃)Cl₂, CH₂=CHCOO(CH₂)₂SiCl₃,
 CH₂=CHCOO(CH₂)₃Si(CH₃)Cl₂, CH₂=CHCOO(CH₂)₃SiCl₃, CH₂=C(CH₃)COO(CH₂)₂Si(CH₃)(OCH₃)₂,
 CH₂=C(CH₃)COO(CH₂)₂Si(OCH₃)₃, CH₂=C(CH₃)COO(CH₂)₃Si(CH₃)(OCH₃)₂, CH₂=C(CH₃)COO(CH₂)₃Si(OCH₃)₃,
 CH₂=C(CH₃)COO(CH₂)₂Si(CH₃)Cl₂, CH₂=C(CH₃)COO(CH₂)₂SiCl₃, CH₂=C(CH₃)COO(CH₂)₃Si(CH₃)Cl₂, and
 CH₂=C(CH₃)COO(CH₂)₃SiCl₃.

5

10



15

20

25

30

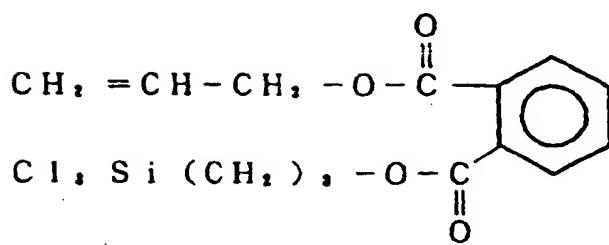
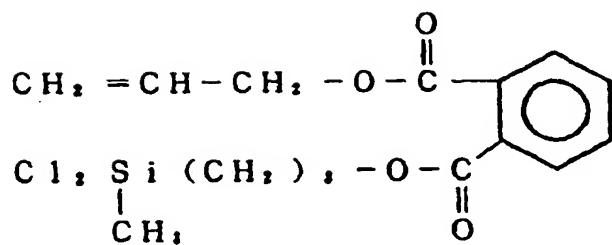
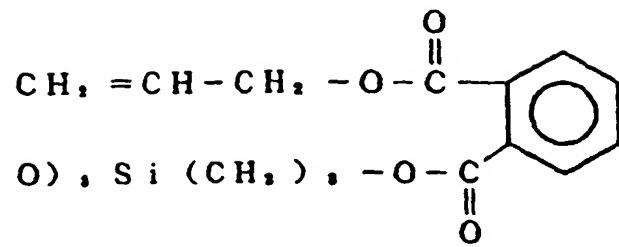
35

40

45

50

55



45 [0087] The compound may be provided in the form of a single compound or a combination of two or more compounds.

[0088] The vinyl monomer to be co-polymerized with the unsaturated silane compound B may be selected from the vinyl monomers having the functional group α previously described with the manner A1 and other similar vinyl monomers.

50 [0089] Alternatively, the component (b) may be selected from specific silyl contained epoxy resins and specific silyl contained polyester resins.

[0090] The specific silyl contained epoxy resin may be produced by causing the epoxy group in an epoxy resin such as bisphenol A type epoxy resin, bisphenol F type epoxy resin, water-added bisphenol A type epoxy resin, aliphatic polyglycidylether, or aliphatic polyglycidylester to react on aminosilane, vinylsilane, carboxysilane, or glycidylsilane having the specific silyl group.

55 [0091] The specific silyl contained polyester resin may be produced by causing the carboxyl group or hydroxy group in a polyester resin to react on aminosilane, carboxysilane, or glycidylsilane having the specific silyl group.

[0092] Mw of the component (b) ranges preferably 2,000 to 100,000 and more preferably 4,000 to 50,000.

[0093] The amount of the component (b) used in the composition (II) is generally 2 to 900 parts by weight, preferably 10 to 400 parts by weight, and more preferably 20 to 200 parts by weight, based on 100 parts by weight of the structural unit $(R^1)_nSiO_{(4-n)/2}$ in the component (a). In that case, when the component is smaller than 2 parts by weight, its resultant coating layer will be declined in the resistance to alkalis. If higher than 900 parts by weight, the resistance to climate of the coating layer throughout a long period of time will be lowered.

[0094] The polymerization of producing the component (b) may be implemented by a process of feeding the monomers at once for polymerization, a process of polymerizing some of the monomers and then continuously or intermittently feeding the remaining monomers, or a process of feeding the monomers in a succession throughout polymerization. Also, any combination of the polymerizing processes may be employed. Most preferable is a technique of solution polymerization. Any common solvent for the solution used in the polymerizing process may be used; a ketone or alcohol solvent is more desirable. For the polymerization, a polymerizing initiator, a molecular weight modifier, a chelating agent, and an inorganic electrolysis may successfully be selected from known agents.

[0095] According to the present invention, the component (b) may be provided in the form of a single substance or a combination of two or more substance as described above.

[0096] Preferably, the component (b) in the composition (II) may be condensed together with the component (a) under the presence of water or an organic solvent.

[0097] The composition (II) of the present invention is composed essentially of the components (a), (b), and (c) and, if desired, may be added with the optional components described later. The organic solvent used for preparing the composition (II) is equally selected from those described previously for the composition (I).

[0098] The component (b) in the composition (II) of the present invention may be a silyl based fluorine polymer (b2) which is described below.

Component (b2)

[0099] The component (b2) is a fluorine polymer having a structural unit expressed by Formula 2 (referred to as a structural unit b2-1 hereinafter) and/or a structural unit expressed by Formula 3 (referred to as a structural unit b2-2 hereinafter) and arranged in which the specific silyl group is preferably linked to an end and/or a side of its polymer molecular chain.

[0100] The component (b2) allows the resultant coating layer made of the composition of the present invention to have improved coating properties because the hydrolytic group and/or the hydroxy group in the specific silyl group is condensed together with the component (a) when the coating layer is being cured.

[0101] The amount of silicon atoms in the component (b2) is generally 0.1 to 60 mol % and preferably 0.5 to 50 mol %, based on the total amount of the component (b2).

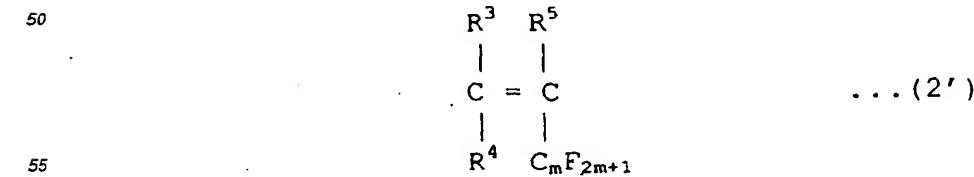
[0102] The component (b2) is preferably synthesized by polymerizing a monomer (referred to as a monomer (b2-1) hereinafter) having the structure unit denoted by Formula 2 and/or a monomer (referred to as a monomer (b2-2) hereinafter) having the structural unit denoted by Formula 3 and a monomer (referred to as a monomer b2-3 hereinafter) having a set of silicon atoms bonded to the hydrolytic group and/or the hydroxy group in the specific silyl group expressed by Formula 6 and may further be added with another monomer (referred to as a monomer (b2-4) hereinafter) which can co-polymerize with the foregoing monomers. The monomer (b2-4) may contain fluorine atoms which are absent in the other monomers (b2-1) and (b2-2).

[0103] The component (b2) may also be a polymer synthesized by polymerizing the monomer (b2-1) and/or the monomer (b2-2); the polymer may be added with the monomer (b2-4), if desired, where its carbon-carbon double bond is linked by reaction to a silane compound (referred to as a silane compound (b2-5) hereinafter) which has a functional group capable of reacting with the hydrolytic group or the hydroxy group.

45

Monomer (b2-1)

[0104] The monomer (b2-1) is expressed by Formula 2'.



(wherein, R^3 to R^5 are $C_m Y_{2m+1}$, m is an integer ranging 0 to 5, and Y is selected from F, H, and Cl separately).

[0105] The monomer (b2-1) may be a compound having a polymeric unsaturated double bond group and at least one fluorine atom.

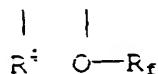
[0106] More particularly, the monomer (b2-1) may be selected from:

5 (A) fluoroethylenes including $\text{CF}_2=\text{CF}_2$, $\text{CHF}=\text{CF}_2$, $\text{CH}_2=\text{CF}_2$, $\text{CH}_2=\text{CHF}$, $\text{CCIF}=\text{CF}_2$, $\text{CHCl}=\text{CF}_2$, $\text{CCl}_2=\text{CF}_2$, $\text{CClF}=\text{CClF}$, $\text{CHF}=\text{CCl}_2$, $\text{CH}_2=\text{CClF}$, and $\text{CCl}_2=\text{CClF}$;
 (B) fluoropropenes including $\text{CF}_3\text{CF}=\text{CF}_2$, $\text{CF}_3\text{CF}=\text{CHF}$, $\text{CF}_3\text{CH}=\text{CF}_2$, $\text{CF}_3\text{CF}=\text{CH}_2$, $\text{CF}_3\text{CF}=\text{CHF}$, $\text{CHF}_2\text{CF}=\text{CHF}$, $\text{CF}_3\text{CH}=\text{CH}_2$, $\text{CH}_3\text{CF}=\text{CF}_2$, $\text{CH}_3\text{CH}=\text{CF}_2$, $\text{CF}_3\text{CF}=\text{CH}_2$, $\text{CF}_3\text{ClCF}=\text{CF}_2$, $\text{CF}_3\text{CCl}=\text{CF}_2$, $\text{CF}_3\text{CF}=\text{CFCI}$, $\text{CF}_3\text{ClCCl}=\text{CCl}_2$, $\text{CCl}_3\text{CF}=\text{CF}_2$, $\text{CF}_3\text{ClCCl}=\text{CCl}_2$, $\text{CFCl}_2\text{CCl}=\text{CHCl}$, $\text{CHF}_2\text{CCl}=\text{CCl}_2$, $\text{CF}_2\text{CICH}=\text{CCl}_2$, $\text{CF}_2\text{CICl}=\text{CHCl}$, and $\text{CCl}_3\text{CF}=\text{CHCl}$; and
 10 (C) Fluoroolephines having not smaller than four carbon atoms including $\text{CF}_3\text{CF}_2\text{CF}=\text{CF}_2$, $\text{CF}_3\text{CF}=\text{CFCF}_3$, $\text{CF}_3\text{CH}=\text{FCF}_3$, $\text{CF}_2=\text{CFCF}_2\text{CHF}_2$, $\text{CF}_3\text{CF}_2\text{CF}=\text{CH}_2$, $\text{CF}_2\text{CH}=\text{CHCF}_3$, $\text{CF}_2=\text{CFCF}_2\text{CH}_3$, $\text{CF}_2=\text{CFCH}_2\text{CH}_3$, $\text{CF}_3\text{CH}_2\text{CH}=\text{CH}_2$, $\text{CF}_3\text{CH}=\text{CHCH}_3$, $\text{CF}_2=\text{CHCH}_2\text{CH}_3$, $\text{CH}_3\text{CF}_2\text{CH}=\text{CH}_2$, $\text{CFH}_2\text{CH}=\text{CHCFH}_2$, $\text{CH}_3\text{CF}_2\text{CH}=\text{CH}_2$, $\text{CF}_2=\text{CFCH}_2\text{CH}_3$, $\text{CF}_3(\text{CF}_2)_2\text{CF}=\text{CF}_2$, and $\text{CF}_3(\text{CF}_2)_2\text{CF}=\text{CF}_2$.

15 [0107] The monomer (b2-1) having fluorine atoms may be provided in the form of a single monomer or a combination of two or more monomers.

Monomer (b2-2)

20 [0108] The monomer (b2-2) is expressed by Formula 3':



35 (wherein, R_f is an alkyl group or an alkoxyalkyl group having fluorine atoms and R^3 to R^5 are analogous to those of Formula 2' but may be modified without departing from the term of analogy to Formula 2').

40 [0109] The monomer (b2-2) may be a compound having a polymeric unsaturated double bond group, an ether bond, and at least one fluorine atom.

[0110] More particularly, the monomer (b2-2) may be selected from:

45 (A) (fluoroalkyl) vinyl ether or (fluoroalkoxyalkyl) vinyl ether expressed by $\text{CH}_2=\text{CH}-\text{O}-\text{R}_f$ (wherein, R_f is an alkyl group or an alkoxyalkyl group having fluorine atoms);
 (B) perfluoro(alkyl)vinyl ethers including perfluoromethylvinyl ether, perfluoroethylvinyl ether, perfluoropropylvinyl ether, perfluorobutylvinyl ether, and perfluoroisobutylvinyl ether; and
 (C) perfluoro(alkoxyalkyl)vinyl ethers including perfluoropropoxypropylvinyl ether.

50 [0111] The monomer (b2-2) having fluorine atoms may be provided in the form of a single monomer or a combination of two or more monomers.

[0112] When the two monomers (b2-1) and (b2-2) are used in a combination, they may preferably be hexafluoropropylene and perfluoroalkyl-perfluorovinyether or perfluoroalkoxyalkyl-perfluorovinyether.

Monomer (b2-3)

55 [0113] The monomer (b2-3) is a monomer having a polymeric unsaturated double bond in one molecule and silicon atoms bonded to a hydrolytic group and/or a hydroxy group.

[0114] The monomer (b2-3) may be an unsaturated silane compound B expressed by Formula 6 used for synthe-

sizing the silyl contained polymer (b).

Monomer (b2-4)

5 [0115] The monomer (b2-4) capable of co-polymerizing with the monomers (b2-1) to (b2-3) may be a vinyl monomer having the functional group α used for synthesizing the silyl contained polymer (b).

Silane compound (b2-5)

10 [0116] The silane compound (b2-5) for addition reaction may be selected from halogenated silanes including methyldichlorsilane, trichlorsilane, and phenyl-dichlorsilane; alkoxy silanes including methyldiethoxysilane, methyldimethoxysilane, phenyl-dimehoxy silane, trimethoxysilane, and triethoxysilane; acyroxysilanes including metylediacetoxysilane, phenyl-diacetoxysilane, and triacetoxysilane; and aminoxy silanes including metyldiaminoxy silane, triaminoxy silane, and dimethylaminoxy silane. The silane compound (b2-5) may be provided in the form of a single compound or a combination of two or more compounds.

15 [0117] The polymerization for synthesizing the component (b2) may be implemented by a process of polymerizing the monomers at once, a process of polymerizing some of the monomers and feeding the remaining of the monomers continuously or intermittently, or a process of feeding the monomers in a succession. Any combination of the foregoing processes may also be used with equal success. Most preferable is a technique of solution polymerization. The solvent 20 used in the solution polymerization is any commonly available solvent and may preferably be selected from ketones and alcohols. For the polymerization, a polymerizing initiator, a molecular weight modifier, a chelating agent, and an inorganic electrolysis may be known ones.

25 [0118] The amount of a sum of the structural units (b2-1) and (b2-2) in the component (b2) is generally 0.5 to 80 mol % and preferably 1 to 70 mol %, based on the total amount of the component (b2). When the amount is lower than 0.5 mol %, the resultant coating layer may hardly be improved in the transparency and the adhesivity. If higher than 80 mol %, the adhesivity of the coating layer to a substrate may be declined.

30 [0119] The amount of the structural unit (b2-1) is preferably 0.5 to 70 mol %, based on the total amount of the component (b2). The amount of the structural unit (b2-2) is preferably 0.5 to 70 mol %, based on the total amount of the component (b2).

35 [0120] The amount of the structural unit (b2-3) is generally 0.1 to 60 mol % of the specific silyl group and preferably 0.5 to 50 mol %, based on the total amount of the component (b2). When the amount is lower than 0.1 mol %, the effect of condensation together with the component (a) will hardly be ensured. If higher than 60 mol %, the storage stability of the resultant coating composition will be declined.

40 [0121] The amount of the structural unit (b2-4) composed of the monomer (b2-4) capable of co-polymerizing with the other monomers is generally not higher than 90 mol % and preferably not higher than 80 mol %, based on the total amount of the component (b2).

[0122] Mw of the component (b2) is preferably 1,000 to 100,000 and more preferably 5,000 to 50,000.

[0123] The component (b2) in the composition (II) may be provided in the form of a single substance or a combination of two or more substance as described above.

45 [0124] The amount of the component (b2) in the composition (II) is generally 20 to 500 parts by weight, preferably 25 to 400 parts by weight, and more preferably 50 to 300 parts by weight, based on 100 parts by weight of the structural unit $(R^1)_nSiO_{(4-n)/2}$ in the component (a). When the amount of component (b2) is smaller than 20 parts by weight, the resultant coating layer will be declined in the resistance to alkalis and cracking. If greater than 500 parts by weight, the resistance to climate of the coating layer will be declined.

[0125] Preferably for the composition (II), the component (b2) is condensed together with the component (a) under the presence of water and/or an organic solvent.

50 [0126] According to the present invention, another vinyl copolymer having the specific silyl group such as the component (b) may be used in addition to the component (b2) without impairing the effect of the present invention. The amount of the another vinyl copolymer in the composition (II) is generally not greater than 500 parts by weight, based on 100 parts by weight of the structural unit $(R^1)_nSiO_{(4-n)/2}$ in the component (a).

[0127] The components (a) and (c) and the organic solvent in the composition (II) are identical in the type and the amount to those in the composition (I) and will be explained in no more detail.

Compositions (i) and (ii)

55 [0128] The composition (i) is similar to the composition (I) while the composition (ii) is similar to the composition (II); the two (i) and (ii) are equivalent to the compositions (I) and (II) respectively less the component (c). The components and the type and amount of an organic solvent in the compositions (i) and (ii) are identical to those in their correspond-

ing compositions (I) and (II) except the component (c) and will be explained in no more detail.

[0129] Each of the compositions (i), (ii), (I), and (II) may further be added with the following components (d) to (h) as well as other additives, which will be explained below. Component (d)

[0130] The composition (i) or (ii) may preferably contain the component (d).

[0131] The component (d) in the composition of the present invention is an ultraviolet light absorber such as fine particles of a semiconductor capable of absorbing ultraviolet light.

[0132] The ultraviolet light absorber may be selected from inorganic semiconductors including TiO_2 in rutile crystalline structure (having no photocatalytic function), ZnO , and CeO_2 ; and organic ultraviolet ray absorbers including salicyl acids, benzophenones, benzotriazoles, cyanoacrylates, and triazines. Most preferable are ZnO and CeO_2 .

[0133] The ultraviolet light absorbing function of the component (d) in the composition of the present invention allows the resultant coating layer to absorb ultraviolet light without trading off the substantial properties, hence preventing deterioration of the undercoating and of the substrate by ultraviolet light.

[0134] The component (d) is provided in three different states: powder of fine particles, aqueous sol where fine particles are dispersed in water, and solvent sol where fine particles are dispersed in a polar solvent such as isopropylalcohol or a non-polar solvent such as toluene. The solvent sol may further be diluted with water or a solvent depending on the degree of dispersion of fine semiconductor particles. The average particle diameter of semiconductor particles in such a state is preferably as small as possible in view of the absorption of ultraviolet light; it is generally not greater than 1 μm , preferably not greater than 0.5 μm , and more preferably not greater than 0.1 μm . Since the component (d) is provided in the form of fine particles, its ultraviolet light absorbing function increases to a level equal to that of an organic ultraviolet light absorber. Also, improvement in the dispersion uniformity results in the higher transparency and the storage stability of the coating. The fine particles or the sol may preferably be added with a surface activator, a dispersing agent, and a coupling agent for conducting further surface treatments.

[0135] When the component (d) is in the form of an aqueous sol or a solvent sol, its solid concentration is preferably not higher than 60 percent by weight and more preferably not higher than 50 percent by weight.

[0136] The method of combining the component (d) with the composition of the present invention may involve either feeding the component (d) after the supply of the other components or feeding the component (d) during preparation of the composition of the present invention and preventing hydrolysis and partial condensation of the organosilanes of the component (a) under the presence of the component (d). Feeding the component (d) during the preparation of the composition allows co-condensation of the semiconductor compounds in the component (d) with the component (a), thus increasing the dispersion property of the component (d). When the component (d) is an aqueous sol, it may preferably be fed during the preparation of the composition. Also, in case that the viscosity in a system is increased by addition of the component (g) described later, the component (d) may preferably be fed during the preparation of the composition.

[0137] The component (d) may commercially be available as Taipak TTO of Ishihara Sangyo, ZW-143 ZW-5130 ZS-300, ZS-303, ZnO -100, and ZnO -200 of Sumitomo Osaka Cement, Z-Nouve of Mitsui Metal Mining, Needlai of Tagi Chemical, and Ceriguard and hicera super K29 of Nihon Inorganic Chemistry.

[0138] The component (d) in the composition of the present invention may be provided in the form of a single substance or a combination of two or more substance.

[0139] The amount of the component (d), based on 100 parts by weight of the structural unit, $(R^1)_nSiO_{(4-n)/2}$, in the component (a) is generally, in solid state, 1 to 150 parts by weight and preferably 5 to 100 parts by weight. When the amount is smaller than 1 part by weight, the ultraviolet light absorbing function will be declined. If greater than 150 parts by weight, the resultant coating layer will be declined in the forming properties, thus suffering from cracking and peeling.

Component (e)

[0140] The compositions (I), (II), (i), and (ii) may contain the component (e).

[0141] According to the present invention, the component (e) is colloidal silica and/or a colloidal alumina. It may be provided in an aqueous sol or colloid where it is dispersed in water, or a solvent sol or colloid where it is dispersed in a polar solvent such as isopropylalcohol or a non-polar solvent such as toluene. The component (e) in the solvent sol or colloid form may further be diluted with water or the solvent depending on the degree of dispersion in the sol or colloid or may be surface treated for increasing the dispersing properties.

[0142] The component (e) of the aqueous sol or colloid, or the solvent sol or colloid form is preferably not higher than 40 percent by weight in the solid concentration.

[0143] The average particle diameter in the colloid form is preferably not greater than 1 μm and more preferably not greater than 0.5 μm . When the average diameter exceeds 1 μm , the particles may sink down.

[0144] The component (e) in the composition of the present invention contributes to the improvement of the flexibility of the resultant coating layer when cured and the increase of its critical film thickness. Also, the component (e) when used for a composition for overcoating (namely (I) or (II)) permits the use of the component (c) to be minimum without affecting the photocatalytic function, thus improving the resistance to climate of the resultant overcoating. Also, the

component (e) when used for a composition for undercoating (namely (i) or (ii)) improves the adhesivity of the overcoating to the undercoating.

[0145] The method of combining the component (e) with the composition may involve feeding the component (e) after the preparation of the composition or feeding the same during the preparation of the composition to allow the hydrolysis and condensation of the component (e) together with the other components (a), (b), and (b2) or their condensates.

[0146] The colloidal silica form of the component (e) may commercially be available as, for example, Snowtex, Iso-propanol Silicagel, and Methanolgel by Nissan Chemical; Cataloid and Oscar by Shokubai Chemical; Ludox by Dupont, U.S.A.; and Nalcoag by Nalco Chemical, U.S.A. The colloidal alumina form of the component (e) maybe available as Alminasol-100, Aluminasol-200, and Aluminasol-520 by Nissan Chemical; and Aluminum Oxide C by Dhexa, West Germany.

[0147] The component (e) in the composition of the present invention may be provided in the form of a single substance or a combination of two or more substances.

[0148] The amount of the component (e) based on 100 parts by weight of the structural unit, $(R^1)_nSiO_{(4-n)/2}$, in the component (a), is generally 0 to 500 parts by weight and preferably 0.1 to 400 parts by weight in solid state.

Component (f)

[0149] At least one of the compositions (I), (II), (i), and (ii) preferably contains the component (f) explained below.

[0150] The component (f) is a catalyst for encouraging the hydrolysis and condensation of the components (a), (b) and (b2).

[0151] The use of the component (f) will accelerate the speed of curing the resultant coating layer and increase the molecular weight of polysiloxane produced by polycondensation of the organosilane component, hence allowing the resultant coating layer to be improved in the physical strength and the long-term durability as well as making the coating layer thickness increased and facilitating the coating application.

[0152] The component (f) is preferably selected from acidic compounds, alkali compounds, basic compounds, amine compounds, metalates, and/or their partial hydrolyzates (the metalates and/or their partial hydrolyzates being referred to as organic metallic compounds).

[0153] Characteristic examples of the acidic compound are acetic acid, hydrochloric acid, sulfuric acid, phosphoric acid, alkyltitanic acid, *p*-toluenesulphonic acid, and phthalic acid and it may preferably be acetic acid.

[0154] Characteristic examples of the alkali compound are sodium hydroxide and potassium hydroxide and it may preferably be sodium hydroxide.

[0155] Characteristic examples of the basic compound are alkali metal salts including naphthenic acid, octyl acid, nitrous acid, sulfuric acid, and aluminate.

[0156] Characteristic examples of the amine compound are ethylene-diamine, hexamethylene-diamine, diethylene-triamine, triethylene-tetramine, tetraethylene-pentamine, piperidine, piperadine, *m*-phenylenediamine, *p*-phenylenediamine, ethanolamine, triethylamine, 3-aminopropyl trimethoxysilane, 3-aminopropyl-triethoxysilane, 3-(2-aminoethyl)-aminopropyl-trimethoxysilane, 3-(2-aminoethyl)-aminopropyl-triethoxysilane, 3-(2-aminoethyl)-aminopropyl-methyl-dimethoxysilane, 3-anilinopropyl-trimethoxysilane, alkylamine salts, class-4 ammonium salts, and modified amines used as a hardener for epoxy resin, and preferably it may be 3-aminopropyl-trimethoxysilane, 3-aminopropyl-triethoxysilane, or 3-(2-aminoethyl)-aminopropyl-trimethoxysilane.

[0157] The organic metallic compound may be selected from a particular compound expressed by Formula 7 shown below (referred to as an organic metallic compound (7) hereinafter), an organic metal compound of quadrivalent tin having 1 to 2 alkyl groups where 1 to 10 carbon atoms are bonded to the corresponding number of tin atoms (referred to as an organic tin compound hereinafter), and their hydrolyzates.



(wherein, M is zirconium, titan, or aluminum, R¹⁰ and R¹¹ are different or identical to each other as, like R⁶, are monovalent hydrocarbon groups having 1 to 6 carbon atoms, R¹² is monovalent a hydrocarbon group having 1 to 6 carbon atoms, like R⁶, or an alkoxy group having 1 to 16 carbon atoms such as methoxy, ethoxy, *n*-propoxy, *i*-propoxy, *n*-butoxy, *sec*-butoxy, *t*-butoxy, lauryloxy, or stearloxy, and r and s are integers ranging from 0 to 4, (r+s)=(valence of M).)

[0158] The organic metallic compound (7) is preferably an alkoxy group having 1 to 16 carbon atoms where r and s are integers ranging from 0 to 4 and (r+s)=(valence of M) being established.

[0159] The organic metallic compound (7) may be selected from:

(A) organic zirconium compounds including tetra-*n*-butoxy zirconium, tri-*n*-butoxy-ethylacetatoacetate zirconium, di-*n*-butoxy-bis(ethylacetatoacetate) zirconium, *n*-butoxy tris(ethylacetatoacetate) zirconium, tetrakis(*n*-propylacetatoacetate) zirconium, and tetrakis(*n*-propylacetatoacetate) titan.

tate) zirconium, tetrakis(acetylacetate) zirconium, and tetrakis(ethylacetate) zirconium;
 (B) organic titan compounds including tetra-i-propoxy titanium, di-i-propoxy bis(ethylacetate) titanium, di-i-propoxy bis(acetylacetate) titanium, and di-i-propoxy bis(acetylacetone) titanium; and
 (C) organic aluminum compounds including tri-i-propoxy aluminum, di-i-propoxy ethylacetate aluminum, di-i-propoxy acetylacetone aluminum, i-propoxy bis(ethylacetate) aluminum, i-propoxy bis(acetylacetone) aluminum, tris(ethylacetate) aluminum, tris(acetylacetone) aluminum, and monoacetylacetone bis(ethylacetate) aluminum.

[0160] The organic metallic compound (7) and its partial hydrolyzate may preferably be selected from tri-n-butoxy ethylacetate zirconium, di-i-propoxy bis(acetylacetone) titanium, di-i-propoxy ethylacetate aluminum, tris(ethylacetate) aluminum, and their partial hydrolyzates.

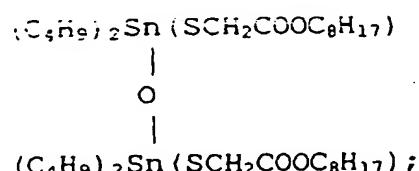
[0161] The organic tin compound may be selected from:

carbonic acid organic tin compounds including $(C_4H_9)_2Sn(OCOC_{11}H_{23})_2$, $(C_4H_9)_2Sn(OCOCH=CHCOOCH_3)_2$,
 $(C_4H_9)_2Sn(OCOCH=CHCOOC_4H_9)_2$, $(C_8H_{17})_2Sn(OCOC_8H_{17})_2$, $(C_8H_{17})_2Sn(OCOC_{11}H_{23})_2$,
 $(C_8H_{17})_2Sn(OCOCH=CHCOOCH_3)_2$, $(C_8H_{17})_2Sn(OCOCH=CHCOOC_4H_9)_2$,
 $(C_8H_{17})_2Sn(OCOCH=CHCOOC_8H_{17})_2$, $(C_8H_{17})_2Sn(OCOCH=CHCOOC_{16}H_{33})_2$,
 $(C_8H_{17})_2Sn(OCOCH=CHCOOC_{17}H_{35})_2$, $(C_8H_{17})_2Sn(OCOCH=CHCOOC_{18}H_{37})_2$,
 $(C_8H_{17})_2Sn(OCOCH=CHCOOC_{20}H_{41})_2$,

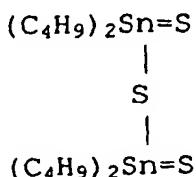


30 $(C_4H_9)Sn(OCOC_{11}H_{23})_2$, and $(C_4H_9)Sn(OCONa)_3$;

35 mercaptide organic tin compounds including $(C_4H_9)_2Sn(SCH_2COOC_8H_{17})_2$, $(C_4H_9)_2Sn(SCH_2CH_2COOC_8H_{17})_2$,
 $(C_8H_{17})_2Sn(SCH_2COOC_8H_{17})_2$, $(C_8H_{17})_2Sn(SCH_2CH_2COOC_8H_{17})_2$, $(C_8H_{17})_2Sn(SCH_2COOC_{12}H_{25})_2$,
 $(C_8H_{17})_2Sn(SCH_2CH_2COOC_{12}H_{25})_2$, $(C_8H_9)_2Sn(SCOCH=CHCOOC_8H_{17})_2$, $(C_8H_{17})_2Sn(SCOCH=CHCOOC_8H_{17})_2$.

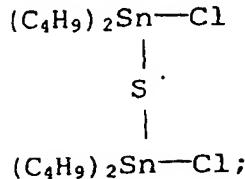


45 sulfide organic tin compounds including $(C_4H_9)Sn=S$, $(C_8H_{17})_2Sn=S$, and



55

chloride organic compounds including $(C_4H_9)SnCl_3$, $(C_4H_9)_2SnCl_2$, $(C_8H_{17})_2SnCl_2$, and



5

10

and

organic tin oxides including $(\text{C}_4\text{H}_9)_2\text{SnO}$ and $(\text{C}_8\text{H}_{17})_2\text{SnO}$ as well as reaction products including an ester compound produced by reaction between the organic oxide and silicate, maleic acid dimethyl, maleic acid diethyl, or phthalic acid diethyl.

[0162] The component (f) may be provided in the form of a single substance or a combination of two or more substances and may be mixed with a zinc compound or a reaction retarder.

[0163] The component (f) may also be fed at a stage for preparation of the composition or at a stage where a coating layer is formed or at both the stages for preparation of the composition and for forming the coating layer.

[0164] The amount of the component (f), except the organic metallic compounds, based on 100 parts by weight of the structural unit, $(\text{R}^1)_n\text{SiO}_{(4-n)/2}$, in the component (a) is generally 0 to 100 parts by weight, preferably 0.01 to 80 parts by weight, and more preferably 0.1 to 50 parts by weight. The amount of the component (f) of any organic metallic compound based on 100 parts by weight of the structural unit, $(\text{R}^1)_n\text{SiO}_{(4-n)/2}$, in the component (a) is generally 0 to 100 parts by weight, preferably 0.1 to 80 parts by weight, and more preferably 0.5 to 50 parts by weight. When the amount of the component (f) exceeds 100 parts by weight, the composition will be declined in the storage stability and its resultant coating layer will suffer from cracking.

Component (g)

[0165] Preferably, at least one of the compositions (i), (ii), (i), and (ii) contains the component (g).

[0166] The component (g) may be at least one expressed by Formula 4 and selected from β -diketones, β -ketesters, carbonic acid compounds, dihydroxy compounds, amine compounds, and oxyaldehydes.

[0167] The component (g) is preferably added when the component (f) is any organic metallic compound.

[0168] The component (g) serves as a stability enhancing agent for the composition. More specifically, it is presumed that the component (g) is coordinates bonded to metallic atoms in the organic metallic compound thus to appropriately control the promotion of co-condensation of the components (a), (b), and (b2) with the organic metallic compound, hence improving the storage stability of the composition.

[0169] Characteristic examples of the component (g) are acetylacetone, acetoacetic acid methyl, acetoacetic acid ethyl, acetoacetic acid-n-propyl, acetoacetic acid-n-propyl, acetoacetic acid-n-butyl, acetoacetic acid-sec-butyl, acetoacetic acid-t-butyl, hexylacetone, and acetoacetic acid ethyl.

[0170] The component (g) may be provided in the form of a single substance or a combination of two or more substances.

[0171] The amount of the component (g) based on 1 mole of the organic metallic compound is generally not smaller than 2 moles and preferably 3 to 20 moles. When the amount of the component (g) is smaller than 2 moles, improvement of the storage stability of the resultant composition will be uncertain.

Component (h)

[0172] The coating composition of the present invention may contain the component (h) described below.

[0173] The component (h) is a powder and/or a sol or a colloid form of an inorganic compound having no photocatalytic function and may be fed so that the resultant coating layer has desired properties (provided that the component (g) is not involved). The sol or colloid form of the component (h) is generally 0.001 to 100 μm in the average particle diameter.

[0174] Characteristic examples of the component (h) are AlGaAs , $\text{Al}(\text{OH})_3$, Sb_2O_5 , Si_3N_4 , $\text{Sn}-\text{In}_2\text{O}_3$, $\text{Sb}-\text{In}_2\text{O}_3$, MgF , CeF_3 , CeO_2 , $3\text{Al}_2\text{O}_3-2\text{SiO}_2$, BeO , SiC , AlN , Fe , Co , $\text{Co}-\text{FeO}_x$, CrO_2 , Fe_4N , BaTiO_3 , $\text{BaO}-\text{Al}_2\text{O}_3-\text{SiO}_2$, Ba ferrite, SmCO_3 , YCO_5 , CeCO_5 , PrCO_5 , $\text{Sm}_2\text{CO}_{17}$, $\text{Nd}_2\text{Fe}_{14}\text{B}$, Al_4O_3 , $\alpha\text{-Si}$, SiN_4 , CoO , $\text{Sb}-\text{SnO}_2$, Sb_2O_5 , MnO_2 , MnB , Co_3O_4 , Co_3B , LiTaO_3 , MgO , MgAl_2O_4 , BeAl_2O_4 , ZrSiO_4 , ZnSb , PbTe , GeSi , FeSi_2 , CrSi_2 , CoSi_2 , $\text{MnSi}_{1.73}$, Mg_2Si , $\beta\text{-B}$, BaC .

BP, TiB_2 , ZrB_2 , HfB_2 , Ru_2Si_3 , TiO_2 (rutile), TiO_3 , $PbTiO_3$, Al_2TiO_5 , Zn_2SiO_4 , Zr_2SiO_4 , $2MgO_2-Al_2O_3-5SiO_2$, Nb_2O_5 , $Li_2O-Al_2O_3-4SiO_2$, Mg ferrite, Ni ferrite, Ni-Zn ferrite, Li ferrite, and Sr ferrite.

[0175] The component (h) may be provided in the form of a single substance or a combination of two or more substances.

[0176] The component (h) may also be provided in a powder form, an aqueous sol or colloidal form where the substance is dispersed in water, or a solvent sol or colloidal form where the substance is dispersed in a polar solvent such as isopropyl alcohol or a non-polar solvent such as toluene.

[0177] The component (h) of the solvent sol or colloidal form may further be diluted with water or the solvent depending on the dispersing properties of the semiconductor substance or may be surface treated for improving the dispersing properties.

[0178] When the component (h) is the aqueous sol or colloidal form, its solid concentration may preferably be not greater than 40 percent by weight.

[0179] The method of combining the component (h) with the composition may involve feeding the component (h) after the preparation of the composition or feeding the same during the preparation of the composition to allows the hydrolysis and condensation together with the components (a), (b), and (b2) or their condensates.

[0180] The amount of the component (h) based on 100 parts by weight of the structural unit, $(R^1)_nSiO_{(4-n)2}$, in the component (a) is generally 0 to 500 parts by weight and preferably 0.1 to 400 parts by weight in solid state. Other additives

[0181] The composition of the present invention may have an appropriate filler added and dispersed for improving the color and the thickness of a resultant coating layer.

[0182] The filler may be selected from non-aqueous organic and inorganic pigments, ceramics, metals, and alloys of a particle, fiber, or scale form, and their oxides, hydroxides, carbides, nitrides, and sulfides.

[0183] Characteristic examples of the filler are iron, copper, aluminum, nickel, silver, zinc, ferrite, carbon black, stainless steel, silicon dioxide, titan oxide for pigment, aluminum oxide, chrome oxide, manganese oxide, iron oxide, zirconium oxide, cobalt oxide, synthetic mullite, aluminum hydroxide, iron hydroxide, silicon carbide, silicon nitride, boron nitride, clay, diatom earth, slaked lime, gypsum, talc, barium carbonate, calcium carbonate, magnesium carbonate, barium sulfate bentonite, mica, zinc green, chrome green, cobalt green, bilidine, Guinier green, cobalt chrome green, shale green, green soil, manganese green, pigment green, ultramarine, deep blue, rocky blue, cobalt blue, cerian blue, copper borate molybdenum blue, copper sulfide, cobalt purple, mars purple, manganese purple, pigment violet, lead suboxide, calcium hydrochloride, zinc yellow, lead sulfide, chrome yellow, yellow soil, cadmium yellow, strontium yellow, titan yellow, litharge, pigment yellow, copper suboxide, cadmium red, serene red, chrome vermillion, Indian red, zinc white, antihuman white, basic lead sulfate, titan white, lithopone, lead silicate, zirconium oxide, tungsten white, lead, zinc white, Bantiscr white, lead phthalate, manganese white, lead sulfate, carbon black, bone black, diamond black, Matmic black, plant black, potassium titanate whisker, and molybdenum disulfide.

[0184] The filler may be provided in the form of a single substance or a combination of two or more substances.

[0185] The amount of the filler based on 100 parts by weight of the solids of the composition is generally not greater than 300 parts by weight.

[0186] Other particular agents to be added to the composition of the present invention, if desired, are known dehydrating agents including orthoformate methyl, orthoacetic acid methyl, and tetraethoxysilane; dispersing agents including polyoxyethylene alkylether, polyoxyethylene alkylphenylether, polyoxyethylene fatty acid ester, polycarbonic acid polymer surfactant, polycarboxylate, polyphosphate, polyacrylate, polyamide ester salt, and polyethylene glycol; thickening agents including cellulose such as methyl cellulose, ethyl cellulose, hydroxyethyl cellulose, hydroxypropyl cellulose, or hydroxypropylmethyl cellulose, castor oil derivative, and ferrosilicate; inorganic foaming agents including ammonium carbonate, ammonium bicarbonate, ammonium subnitrate, sodium borohydride, and calcium azide; organic foaming agents including an azo compound such as azo-bisisobutyro-nitrile, a hydrazine compound such as diphenylsulfone-3,3'-disulfonyldiazine, a semicarbazide compound, a triazole compound, and an N-nitroso compound; and other additives including a surfactant, a silane coupling agent, a titan coupling agent, and a dye.

[0187] Also, the composition (i) or (ii) for undercoating may be added with an ultraviolet light stabilizer for improving the resistance to climate and the long-term adhesivity. The ultraviolet light stabilizer may be a piperidine.

[0188] Moreover, a leveling agent may be combined for improving the coating performance of the composition. The leveling agent may commercially be available: as a fluorine leveling agent, BM1000 and BM1100 by BM-Chemie, Efca 772 and Efca 777 by Efca Chemicals, a Florence series by Kyoeisha Chemical, an FC series by Sumitomo 3M, and a Fluonal TF series by Toho Chemical; as a silicone leveling agent, a BYK series by BYK Chemie, a Sshmegro series by Sshmegmann, and Efca 30, Efca 31, Efca 34, Efca 35, Efca 36, Efca 39, Efca 83, Efca 86, and Efca 88 by Efca Chemicals; and as an ether or ester leveling agent, Carphynol by Nisshin Chemical and Emargen and Homogenol by Kaoh.

[0189] The use of the leveling agent will improve the appearance of a finished coating layer and also allow the coating layers to be coated uniformly as a thin film.

[0190] The amount of the leveling agent to be used, based on the entirety of the component, is preferably 0.01 to 5

percent by weight and more preferably 0.02 to 3 percent by weight.

[0191] The method of combining the leveling agent may involve feeding at a stage for preparing the composition or at a stage for forming a coating layer or at both the stages for preparing the composition and for forming the coating layer.

[0192] The composition of the present invention or more particularly, the composition (i) or (ii) for undercoating may be blended with another resin. Characteristic examples of the another resin are acryl-urethane resin, epoxy resin, polyester, acryl resin, fluorine resin, acryl resin emulsion, epoxy resin emulsion, urethane emulsion, and polyester emulsion.

[0193] The preparation of the composition of the present invention is not limited to a specific mixing method when the two components (f) and (g) are unused. In case that the components (f) and (g) are used, a process may preferably be employed of preparing a mixture of the components (a) to (h) excluding the component (g) and then doping the component (g) into the mixture.

[0194] The concentration of all solids in the composition (I) or (II) of the present invention is generally 3 to 50 percent by weight and preferably 5 to 40 percent by weight and may be adjusted corresponding to the purpose of application. When the solid concentration in the composition exceeds 50 percent by weight, the storage stability will be declined.

[0195] The concentration of all solids in the composition (i) or (ii) of the present invention is generally not higher than 50 percent by weight and preferably not higher than 40 percent by weight and may be adjusted depending on the type of a substrate, the method of coating, the thickness of a coating layer, and so forth.

20

Photocatalyst Coating Film

[0196] The photocatalyst coating film of the present invention consists mainly of ① a substrate/a composition (i) or (ii)/a composition (I) or (II), or ② a substrate/a primer/a composition (i) or (ii)/a composition (I) or (II).

[0197] The application of any of the compositions of the present invention may be conducted by a known manner such as dip coating, flow coating, spraying, screening, electric deposition, or vapor deposition or with the use of a brush, a roll coater, a flow coater, a centrifugal coater, an ultrasonic coater, or a (micro) gravure coater.

[0198] In application of the composition (i) or (ii) of the present invention, the thickness of a coating layer is substantially 0.05 to 20 μm , when dried, with one single application and 0.1 to 40 μm with two times of application. After dried at normal temperatures or heated to a temperature of 30 to 200°C commonly for 0.5 to 60 minutes the coating layer will set on the substrate of a desired type.

[0199] In application of the composition (I) or (II) of the present invention, the thickness of a coating layer is substantially 0.05 to 20 μm , when dried, with one single application and 0.1 to 40 μm with two times of application. After dried at normal temperatures or heated to a temperature of 30 to 200°C commonly for 0.5 to 60 minutes, the coating layer will set on the substrate of a desired type.

[0200] The total thickness of the undercoating and the overcoating may normally be 0.1 to 80 μm and preferably 0.2 to 60 μm when dried.

Substrate

[0201] The substrate which is able to associate with the composition of the present invention is provided as an organic film selected from polyesters including polyethylene terephthalate (PET), polybutylene terephthalate (PBT), and polyethylene-2,6-naphthalate (PEN); polyamides including nylon 6 and nylon 6,6; polyolefins including polyethylene (PE) and polypropylene (PP); and polyacryls including polycarbonates (PC) and polymethacrylate methyl (PMMA).

[0202] The organic film may be a fluorine film such as polytetrafluoroethylene (PTFE) or ethylene-tetrafluoroethylene (ETFE).

[0203] Characteristic examples of the surface treatment are blasting, chemical treatment, degreasing, flaming, oxidation, steam treatment, corona discharge, ultraviolet radiation, plasma treatment, and ion treatment.

[0204] The method of applying the composition of the present invention may be varied depending on the type and the condition of the substrate as well as the technique to be used. For example, a primer may also be used or not corresponding to the use.

[0205] The primer is not limited to a specific type and may be selected from a variety of materials which enable to enhance the bonding between the substrate and the composition, according to the type of the substrate and the purpose of use. The primer may also be provided in the form of a single substance or a combination of two or more substances. The primer may be an enamel containing a coloring material such as a pigment or may have transparency without such a coloring material.

[0206] Characteristic examples of the primer are alkyd resin, aminoalkyd resin, epoxy resin, polyester, acryl resin,

urethane resin, fluorine resin, acrylsilicone resin, acryl resin emulsion, epoxy resin emulsion, polyurethane emulsion, and polyester emulsion. The primer may have various functional radicals when a higher degree of adhesivity is required between the film substrate and the coating layer in hostile conditions. The functional radicals include, for example, a hydroxy group, a carboxyl group, an amide group, an amine group, a glycidyl group, an alkoxysilyl group, an ether bond, and an ester bond. The primer may also contain an ultraviolet light absorber, an ultraviolet light stabilizer, and so forth.

5 [0207] For increasing the wear resistance and the glossiness, the coating layer of the composition of the present invention may be protected at the upper surface with a clear layer composed of, for example, a siloxane resin paint which may be a stable dispersion solution of colloidal silica and siloxane resin such as disclosed in US Patent Nos. 3,986,997 and 4,027,073.

10 [0208] As set forth above, the photocatalyst coating film according to the present invention is fabricated by providing a coating layer of the composition (i) or (ii) on the upper surface of an organic film substrate and providing a coating layer of the composition (I) or (II) on the coating layer of the composition (i) or (ii). The resultant photocatalyst coating film is improved in the resistance to fouling, the durability, and the transparency and has a photocatalytic function, hence preventing the (organic film) substrate from being deteriorated by exposed to light and ensuring the long-term durability 15 and adhesivity.

Embodiment

20 [0209] The embodiment of the present invention will be described in more detail referring to some examples. It would be understood that the present invention is not limited to the following examples.

[0210] The amounts in the examples and their relevant preparations are denoted in the unit of parts or percent on a weight basis, unless otherwise specified. The measurement and estimation of each characteristic factor was carried out by the following manners.

25 (1) Mw

[0211] A gel permeation chromatography (GPC) method was used under the following conditions.

30 Test item: Tetrahydrofuran was used as a solvent. 1 g of a partial condensate of organosilane or 0.1 g of a silyl contained vinyl resin was dissolved in 100 cc of the tetrahydrofuran to prepare a test item.

Standard polystyrene: Standard polystyrene used was made by Pressure Chemical, U.S.A.

Apparatus: A high-temperature, high-speed gel permeation chromatogram (a model, 150-C ALC/GPC) by Waters, U.S.A.

Column: A model, Shodex A-80M (length 50cm) by Showa Denko, Japan.

55 Measurement temperature: 40 °C.

Flow velocity: 1 cc/min.

(2) Adhesion

40 [0212] A tape peeling test was conducted three times using a matrix (of 100) specified in JIS K5400 and its average was measured.

(3) Hardness

45 [0213] This measurement was based on a pencil hardness specified by JIS K4500.

(4) Resistance to alkalis

50 [0214] After a coating test piece was immersed in a saturated calcium hydroxide solution for 60 days, its coating layers were examined by visual check. The test piece exhibiting no change was classified as "good".

(5) Resistance to organic chemicals

55 [0215] The coating test piece was applied with 2cc of isopropyl alcohol, and wiped off after 5 minutes. Then, the test piece was visually examined. The test piece exhibiting no change was classified as "good".

(6) Resistance to moisture

[0216] After a test piece was left at a temperature of 50 °C and a moisture of 95 % for 1,000 hours continuously, its coating layers were visually examined. The test piece exhibiting no change was classified as "good".

5

(7) Resistance to climate

[0217] A 3000-hour irradiation test with a Sunshine weather meter made by Suga Test was carried out conforming to JIS K5400 and the coating layers of a test piece were visually examined for the state of appearance (cracking and peeling). The test piece exhibiting no change was classified as "good".

10 (8) Resistance to water

[0218] After a test piece was immersed in tap water at room temperature, the coating layers of a test piece were visually examined for the state of appearance (cracking and peeling). The test piece exhibiting no change was classified as "good".

15 (9) Resistance to fouling

[0219] After the coating layers of a test piece were fouled with a mixture paste of carbon black/kerosine oil=1/2 (in weight ratio), left at room temperatures for 24 hours, and rinsed with water using a sponge, it was visually examined. The evaluation was based on the following criteria.

20 ○: Not fouled.

25 Δ: Slightly fouled.

X: Terribly fouled.

30 (10) Transparency

[0220]

35 ○: Transparency of higher than 80 %.

○: Transparency between 60 and 80 %.

Δ: Transparency of smaller than 60 %.

40 Provision 1 (of the component (b))

[0221] In a reactor equipped with a circulating cooler and a stirrer, 70 parts of methylmethacrylate, 40 parts of n-butylacrylate, 20 parts of γ -methacryloxypropyltrimethoxysilane, 5 parts of acrylic acid, 13 parts of 2-hydroxyethylmethacrylate, 1 part of 1,1,1-trimethylamine-methacrylimide, 1 part of 4-(meth)acryloyloxy-2,2,6,6-tetramethylpiperidine, 150 parts of i-propyl alcohol, 50 parts of methylethylketone, and 25 parts of methanol were mixed together. The mixture was then heated to 80 °C while being stirred and applied throughout a period of 30 minutes with drops of a solution prepared by dissolving 4 parts of azobisisovaleronitril into 10 parts of xylene. After the reaction at 80 °C for 5 hours, a polymer solution (referred to as (b-A) hereinafter) having a solid concentration of 40 % and an Mw of 11,000 was synthesized.

45 50 Provision 2 (of the component (b))

[0222] In a reactor equipped with a circulating cooler and a stirrer, 70 parts of methylmethacrylate, 40 parts of n-butylacrylate, 20 parts of γ -methacryloxypropyl-trimethoxysilane, 18 parts of glycidylmethacrylate, 1 part of 1,1,1-trimethylamine-methacrylimide, 1 part of 4-(meth)acryloyloxy-2,2,6,6-tetramethylpiperidine, 150 parts of i-propyl alcohol, 50 parts of methylethylketone, and 25 parts of methanol were mixed together. The mixture was then heated to 80 °C while being stirred and applied throughout a period of 30 minutes with drops of a solution prepared by dissolving 4 parts of azobisisovaleronitril into 10 parts of xylene. After the reaction at 80 °C for 5 hours, a polymer solution (referred to as (b-B) hereinafter) having a solid concentration of 40 % and an Mw of 13,000 was synthesized.

Provision 3 (of the component (b))

[0223] In a reactor equipped with a circulating cooler and a stirrer, 70 parts of methylmethacrylate, 40 parts of n-butyleacrylate, 20 parts of γ -methacryloxypropyl-trimethoxysilane, 18 parts of hydroxyacrylate, 10 parts of acrylamide, 1 part of 1,1,1-trimethylamine-methacrylimide, 1 part of 4-(meth)acryloxy-2,2,6,6-tetramethylpiperidine, 150 parts of i-propyl alcohol, 50 parts of methylethylketone, and 25 parts of methanol were mixed together. The mixture was then heated to 80 °C while being stirred and applied throughout a period of 30 minutes with drops of a solution prepared by dissolving 4 parts of azobisisovaleronitril into 10 parts of xylene. After the reaction at 80 °C for 5 hours, a polymer solution (referred to as (b-C) hereinafter) having a solid concentration of 40 % and an Mw of 10,000 was synthesized.

10 Provision 4 (of the component (b2))

[0224] A stainless autoclave equipped with an electromagnetic stirrer was subjected to substitution with a nitrogen gas and filled with 250 parts of methylisobutylketone, 30 parts of ethylvinylether, and 2 parts of lauroyl peroxide (a radical polymer initiator) to prepare a solution. The solution was then cooled down to -50 °C with dry ice-methanol and oxygen in the autoclave was removed out using a nitrogen gas. Then, the remaining was added with 20 parts of hexafluoropropylene, 45 parts of perfluoro(methylvinylether), and 5 parts of vinyl trimethoxysilane and heated up. When the temperature in the autoclave was increased up to 60 °C, the interior pressure was 5 kgf/cm².

[0225] The mixture was stirred while being kept at a reacting temperature of 60 °C to continue the reaction for polymerization for 20 hours. When the pressure in the autoclave dropped down to 1.5 kgf/cm², the mixture was cooled down with water to cease the reaction. A resultant polymer solution was 40 % in the solid concentration and 30,000 in the Mw (referred to as (b2-A)). Provision 5 to 7 (of the component b2).

[0226] By the same manner as of Provision 4 except that the monomers used were as shown in Table 1, polymer solutions (b2-B to b2-D) having a solid concentration of 40 % were prepared.

25

Table 1

Provisions	4	5	6	7
Polymer	B2-A	B2-B	B2-C	B2-D
Monomer (b2-1), hexafluoropropylene	20	-	65	10
Monomer (b2-2), perfluoro-Methylvinylether	45	65	-	10
Monomer (b2-3), vinyltrimethoxysilane	5	5	5	50
Monomer (b2-4), ethylenevinylether Solvent, methylisobutylketone	30	30	30	30
Mw	30,000	30,000	30,000	25,000

40

Provision 8 (of a primer)

[0227] 100 parts of the polymer solution (b2-A) of Preparation 1 was mixed and stirred with 80 parts of methylethylketone and 20 parts of i-propyl alcohol. Then, the mixture was added with 20 parts of a zinc oxide (having 30 % of a dispersed-in-toluene solid concentration) adjusted by methylethylketone to a solid concentration of 20 % thus to prepare a primer having 20 % of a solid concentration (referred to as (p-1) hereinafter).

Preparations 1 to 27 (of the composition for overcoating)

50 [0228] In a reactor equipped with a stirrer and a circulating cooler, a combination of the components listed in Tables 2 to 7 (excluding acetylacetone) were mixed up and stirred for reaction at 60 °C for 4 hours. Then, the mixture was added with an acetylacetone and cooled down to a room temperature to prepare a composition having a solid concentration of 20 %. 100 parts of the composition was mixed and stirred with 100 parts of i-propyl alcohol and further added with 10 parts of an i-propyl alcohol solution (15 % of a solid) of dibutyl-tin-acetate and a silicate oligomer to yield the 55 overcoating compositions (I-1) to (I-9) and (II-1) to (II-18).

Table 2

Preparations	1	2	3	4	5
Compositions	I-1	I-2	I-3	I-4	I-5
Mixture (in parts)					
Component (a)					
tetraethoxysilane		30	30		
methylsilicate (MS51)				30	
methyltrimethoxysilane	70	70	40	70	70
methylsilicone resin ¹					30
dimethyldimethoxysilane	30		30		
Component (c)					
TiO ₂ dispersed in water ²	250	250	250	250	250
Component (f)					
di-i-propoxy-ethyl-acetoacetatealuminum hydrochloric acid (10 ⁻² mol/l)	10	10	10	10	10
Solvent					
Water	(200)	(200)	(200)	(200)	(200)
i-propyl alcohol	70	70	70	70	70
diethyleneglycol-monoethyl ether	70	70	70	70	70
Additive (in parts)					
Component (g)					
acetylacetone	10	10	10	10	10

35

40

45

50

55

Table 3

Preparations	6	7	8	9
Compositions	I-6	I-7	I-8	I-9
Mixture (in parts)				
Component (a)				
tetraethoxysilane				
methylsilicate (MS51)				
methytrimethoxysilane	70	70	70	70
methylsilicon resin ^{*1}				
dimethyldimethoxysilane		30	30	30
polydimethylsiloxane ^{*2}	30			
Component (c)				
TiO ₂ dispersed in water ^{*3}	250		850	40
TiO ₂ dispersed in IPA ^{*4}		250		
Component (f)				
di-i-propoxy-ethyl-acetoacetatealuminum	10		10	10
hydrochloric acid (10 ⁻² mol/l)		30		
Solvent				
Water	(200)		(680)	(32)
i-propyl alcohol	70	55	70	70
diethyleneglycol-monoethylether	70	55	70	70
Additive (in parts)				
Component (g)				
acetylacetone	10	10	10	10

Table 4

Preparations	10	11	12	13	14
Compositions	II-1	II-2	II-3	II-4	II-5
Mixture (in parts)					
Component (a)					
methylsilicate (MS51)			40		30
methytrimethoxysilane	70	60	60	70	40
methylsilicone resin ¹		40			
dimethyldimethoxysilane	30				30
polydimethylsiloxane ²				30	
Component (b)					
(b-A) having 40% solids	50	50	50	50	
(b-B) having 40% solids					50
Component (c)					
TiO ₂ dispersed in water ³	350	350	350	350	350
TiO ₂ dispersed in IPA ⁴					
Component (f)					
di-i-propoxy-ethyl-acetoacetatealuminum	10	10	10	10	10
water	(280)	(280)	(280)	(280)	(280)
i-isopropyl alcohol	100	100	100	100	100
diethyleneglycol-monoethylether	100	100	100	100	100
Additive (in parts)					
Component (g)					
acetylacetone	10	10	10	10	10

5

10

15

20

25

30

35

40

45

50

55

Table 5

Preparations	15	16	17	18
Compositions	II-6	II-7	II-8	II-9
Mixture (in parts)				
Component (a)				
tetraethoxysilane				
methylsilicate (MS51)				
methytrimethoxysilane	70	70	70	70
methylsilicone resin ¹				
dimethyldimethoxysilane	30	30	30	30
polydimethylsiloxane ²				
Component (b)				
(b-A) having 40% solids		50	50	120
(b-B) having 40% solids	50			
Component (c)				
TiO ₂ dispersed in water ³		850	40	350
TiO ₂ dispersed in IPA ⁴	350			
Component (f)				
di-i-propoxy ethyl-acetoacetatealuminum	10	10	10	10
Water	30	(680)	(32)	(280)
i-propyl alcohol	85	100	90	100
diethylene glycol-monoethyl ether	85	100	90	100
Additive (in parts)				
Component (g)				
acetylacetone	10	10	10	10

40

45

50

55

Table 6

Preparations	19	20	21	22	23
Compositions	II-10	II-11	II-12	II-13	II-14
Mixture (in parts)					
Component (a)					
methytrimethoxysilane	70	70	70	70	70
dimethyldimethoxysilane	30	30	30	30	30
Component (b)					
(b2-A) having 40% solids	50				
(b2-B) having 40% solids		50			
(b2-C) having 40% solids			50		
(b2-D) having 40% solids				50	30
(b-A) having 40% solids					20
Component (c)					
TiO ₂ dispersed in water ⁵	100	100	100	100	100
TiO ₂ dispersed in toluene ⁶					
Component (f)					
tris(ethylacetacetate)-aluminum	5	5	5	5	5
water	(70)	(70)	(70)	(70)	(70)
methyl ethyl ketone	200	200	200	200	200
i-isopropyl alcohol	50	50	50	50	50
Additive (in parts)					
Component (g)					
acetylacetone	5	5	5	5	5

5

10

15

20

25

30

35

40

45

50

55

Table 7

Preparations	24	25	26	27
Compositions	II-15	II-16	II-17	II-18
Mixture (in parts)				
Component (a)				
methytrimethoxysilane	70	70	70	70
dimethyldimethoxysilane	30	30	30	30
Component (b2)				
(b2-A) having 40% solids	50	50	50	50
Component (c)				
TiO ₂ dispersed in water ⁵	12	50		100
ZnO dispersed in toluene ⁶			50	
Component (f)				
tris(ethylaceto-acetatealuminum	5	5	5	5
Component (e)				
SiO ₂ dispersed in MEK ⁷				
Water	(8)	(35)	10	(70)
Methylethylketone	160	185	160	200
i-propyl alcohol	40	45	40	50
Additive (in parts)				
Component (g)				
acetylacetone	5	5	5	5

[0229] Throughout Tables 2 to 7, *1 to *7 are as follows,

- *1: Methylsilicone resin (Mw:500) made by Shinetsu Chemical,
- *2: Polydimethylsiloxane containing ethoxysilane (Mw:5,000) Mw made by Nippon Unica,
- *3: Anatase type titan oxide (20% concentration in solid) dispersed in water (pH4),
- *4: Anatase type titan oxide (20% concentration in solid) dispersed in i-propylalcohol,
- *5: Anatase type titan oxide (30% concentration in solid) dispersed in water (pH4),
- *6: Zinc oxide (30% concentration in solid) dispersed in toluene,
- *7: Silica sol (30% concentration in solid) dispersed in methylethylketone.

Preparation 28 (of the composition for overcoating)

[0230] In a reactor equipped with a stirrer and a circulating cooler, a combination of the components (excluding the additive component) listed in Table 8 were mixed up and stirred for reaction at 60 °C for 4 hours. Then, the mixture was added with 5 parts of acetylacetone, stirred for one hour, and cooled down at a room temperature. The mixture was further added with i-propyl alcohol, diethylglycol monoethylether, and methanol of the additive component to yield the overcoating composition (I-10) having a solid concentration of 10 %.

Preparations 29 and 43 (of the composition for overcoating)

[0231] In a reactor equipped with a stirrer and a circulating cooler, a combination of the components (excluding the component (c), the water, and the additive component) listed in Table 8 or 10 were mixed up and heated to 50 °C while being stirred. Then, the mixture was added with 30 parts of water by drop application throughout a period of 30 minutes

EP 1 022 318 A2

and left for reaction at 60 °C for 4 hours. The mixture was then added with 5 parts of acetylacetone, stirred for one hour, and cooled down to a room temperature. The mixture was further added with the component (c) and the additive component containing i-propyl alcohol, diethylglycol monoethylether, and methanol listed in Table 8 or 10 to yield the overcoating compositions (I-11) and (II-22) having a solid concentration of 10 %.

5 Preparations 30 to 39 and 44 to 46 (of the composition for overcoating)

[0232] In a reactor equipped with a stirrer and a circulating cooler, a combination of the components (excluding the component (c), the component (e), the water, and the additive component) listed in Table 8 or 10 were mixed up and heated to 50 °C while being stirred. Then, the mixture was added with 30 parts of water by drop application throughout a period of 30 minutes and left for reaction at 60 °C for 4 hours. The mixture was then added with 5 parts of acetylacetone, stirred for one hour, and cooled down to a room temperature. The mixture was further added with the component (c), the component (e), and the additive component containing i-propyl alcohol, diethylglycol monoethylether, and methanol listed in Table 8 or 10 to yield the overcoating compositions (I-12) to (I-21) and (II-23) to (II-25) having a solid concentration of 10 %.

10 Preparations 40 to 42 (of the composition for overcoating)

[0233] In a reactor equipped with a stirrer and a circulating cooler, a combination of the components (excluding the component (e) and the additive component) listed in Table 11 were mixed up and stirred for reaction at 60 °C for 4 hours. Then, the mixture was added with 6 parts of acetylacetone, stirred for one hour, and cooled down to a room temperature. The mixture was further added with the component (e) and the additive component containing i-propyl alcohol, diethylglycol monoethylether, and methanol listed in Table 11 to yield the overcoating compositions (II-19) to (II-21).

25

30

35

40

45

50

55

Table 8

Preparations	28	29	30	31	32	33
Compositions	I-10	I-11	I-12	I-13	I-14	I-15
Mixture (in parts)						
Component (a)						
methytrimethoxysilane	101	101	81	81	122	122
dimethyldimethoxysilane				16	16	16
3-glycidydoxypropyl-			14			
trimethoxysilane						
(a2-1)	50	50				
(a2-2)			50			
(a2-3)				50		
(a2-4)					30	30
Component (c)						
TiO ₂ dispersed in water (20%)	300	150	150	200	300	150
Component (e)						
colloidal silica (30% methanol dispersion)		100				100
colloidal alumina (20% water dispersion)			150			
water		30	30	30	30	30
i-propylalcohol	82	152	158	156	135	135
Component (f)						
di-propoxyethyl-acetoacetatealuminum	5	5	5	5	5	5
Additive (in parts)						
Component (g)						
acetylacetone	5	5	5	5	5	5
i-propylalcohol	355	339	322	289	322	339
diethyleneglycol-monoethylether	355	339	322	289	322	339
methanol	355	339	322	289	322	339

Table 9

Preparations	34	35	36	37	38	39
Compositions	I-16	I-17	I-18	I-19	I-20	I-21
Mixture (in parts)						
Component (a)						
methyltrimethoxysilane	122	122	122	81	81	81
dimethylmethoxysilane						
3-glycidoxypropyl-	14	14		14	14	14
trimethoxysilane						
(a2-1)				20	20	20
(a2-2)						
(a2-3)						
(a2-4)	30	30	40	30	30	30
Component (c)						
TiO ₂ dispersed in water (20%)	300	200	150	300	200	150
Component (e)						
colloidal silica (30% methanol dispersion)			100			100
colloidal alumina (20% water dispersion)						
water	30	30	30	30	30	30
i-propylalcohol	137	137	141	158	158	158
Component (f)						
di-i-propoxy-ethyl-acetoacetatealuminum	5	5	5	5	5	5
Additive (in parts)						
Component (g)						
acetylacetone	5	5	5	5	5	5
i-propylalcohol	322	289	339	322	289	339
diethyleneglycol-monoethylether	322	289	339	322	289	339
methanol	322	289	339	322	289	339

45 (wherein, (a2-1) is a trifunctional oligomer (Mw:800) containing terminal alkoxysilyl group, (a2-2) is a trifunctional oligomer (Mw: 8,000) containing terminal alkoxysilyl group, (a2-3) is a trifunctional oligomer (Mw:35,000) containing terminal alkoxysilyl group, and (a2-4) is a bifunctional oligomer (Mw:10,000) containing terminal alkoxysilyl group/poly(oxyethylene/ oxypropylene).)

Table 10

Preparation	40	41	42	43
Compositions	II-19	II-20	II-21	II-22
Mixture (in parts)				
Component (a)				
methytrimethoxysilane	81	81	101	101
dimethyldimethoxysilane	49	32	49	32
γ -methacryloxypropyl-trimethoxysilane			3	
Component (b)				
(b-A) having 40% solids	75	100		
(b-B) having 40% solids			50	75
Component (c)				
colloidal silica (30% methanol dispersion)	67	100		133
colloidal silica (20% water dispersion)			100	
Component (d)				
TiO ₂ dispersed in water (20%)	100	100	150	50
Water	58	56	56	33
i-propyl alcohol	58	56	56	33
diethyleneglycol-monoethylether	58	56	56	33
methanol				
Component (f)				
di-i-propoxy-ethyl-acetacetate-aluminum	6	6	6	6
Additive (in parts)				
Component (g)				
acetylacetone	6	6	6	6
i-propyl alcohol	285	306	283	327
diethyleneglycol-monoethylether	285	306	283	327
methanol	285	306	283	327

Table 11

Preparation	44	45	46
Compositions	II-23	II-24	II-25
Mixture (in parts)			
Component (a)			
methytrimethoxysilane	101	61	61
dimethyldimethoxysilane	32	49	49
γ -glycidyldoxypropyl-trimethoxysilane		14	14
γ -methacryloxypropyl-trimethoxysilane	2	2	2
Component (b)			
(b-A) having 40% solids	75	75	
(b-B) having 40% solids			75
Component (c)			
colloidal silica (30% methanol dispersion)	67	67	100
colloidal alumina (20% water dispersion)			
Component (d)			
TiO ₂ dispersed in water (20%)	100	100	100
Water	25	25	25
Component (f)			
di-i-propoxy-ethyl-acetoacetatealuminum	6	6	6
Additive (in parts)			
Component (g)			
acetylacetone	6	6	6
i-propyl alcohol	333	336	359
diethyleneglycol-monoethylether	333	336	359
methanol	333	336	359

Preparations 47 to 51 (of the composition for undercoating)

[0234] In reactor equipped with a circulating cooler, a combination of the components (excluding the water and the additive component) as listed in Table 12 were mixed together, stirred, and added with 30 parts of water as being stirred for reaction at 60°C for 4 hours. Then, the mixture was combined with the additive component and cooled down to a room temperature to prepare a composition having 30% of a solid concentration. 100 parts of the composition was mixed with 100 parts of i-butyl alcohol and 100 parts of propyleneglycol monomethyletheracetate and further added with 10 parts of an i-propylalcohol solution (15% solids) of dioctyl-tin-dimaleate-ester to yield the undercoating compositions (i-1), (i-2), and (ii-1) to (ii-3).

Preparations 52 to 57 (of the composition for undercoating)

[0235] In reactor equipped with a circulating cooler, a combination of the components (excluding the component (e), the ultraviolet light absorber, the water, and the additive component) as listed in Table 13 were mixed and heated up to 50 °C while being stirred. Then, the mixture was added with 20 parts of water throughout a period of 30 minutes and left for reaction at 60 °C for 4 hours. Then, the mixture was combined with acetylacetone, stirred for one hour, and cooled down to a room temperature. The mixture was further added with the component (e), the ultraviolet light absorber, and the additive component to yield the undercoating compositions (ii-4) to (ii-9).

Table 12

Preparations	47	48	49	50	51
Compositions	i-1	i-2	ii-1	ii-2	ii-3
Mixture (in parts)					
Component (a)					
methytrimethoxysilane	70	70	70	70	70
dimethyldimethoxysilane	30	30	30	30	30
Component (b)					
(b-A) having 40% solids			100	100	80
(b2-A) having 40% solids					20
Component (d)					
ZnO dispersed in IPA ¹⁸		15		15	15
Component (f)					
di-i-propoxy-ethyl-acetoacetatealuminum	10	10	10	10	10
Solvent					
water	30	30	30	30	30
i-propylalcohol	30	30	30	30	30
ethyleneglycol-monobutylether	30	30	30	30	30
Additive (in parts)					
Component (g)					
acetylacetone	10	10	10	10	10

¹⁸ Zinc oxide dispersed in i-propylalcohol (30% concentration in solid)

35

40

45

50

55

Table 13

Preparations	52	53	54	55	56	57
Compositions	ii-4	ii-5	ii-6	ii-7	ii-8	ii-9
Mixture (in parts)						
Component (a)						
methyltrimethoxysilane	41	41	71	71	71	58
dimethyldimethoxysilane	16	16	24	24	24	19
Component (b)						
(b-A) having 40% solids		175				150
(b-B) having 40% solids	175		125	125	125	
Component (d)						
ZnO ₂ (IPA dispersion, 10% solids)	200	200				
CeO ₂ (IBA dispersion, 10% solids)			300			
2,4-hydroxybenzophenone				5		
2-(2'-hydroxy-5'-methylphenyl)benzotriazole				15		
Component (e)						
colloidal silica (IPA dispersion, 10%)			300	300	300	300
water	20	20	20	20	20	20
i-propylalcohol	54	54	62	62	62	58
methylethylketone	27	27	31	31	31	29
Component (f)						
di-i-propoxyethyl-acetoacetatealuminum	3	3	3	3	3	3
Additive (in parts)						
Component (g)						
acetylacetone	3	3	3	3	3	3
i-butylalcohol	166	166	222	222	222	166
metylisobutylketone	417	417	444	444	444	417
ethyleneglycol-monobutylether	83	83				83

Examples 1 to 32

[0236] Each of the undercoating compositions listed in Table 12 was applied to the surface of a 50 μm -thick PET film to have an undercoating of 1 μm in thickness when dried. Then, a corresponding type of the overcoating composition was applied to the undercoating to have an overcoating of 0.2 μm in thickness when dried. After cured, a resultant coating film was analyzed for a variety of the characteristics.

[0237] The result of the analysis is shown in Tables 14 to 17.

Table 14

Examples	1	2	3	4	5	6	7	8
Undercoating	i-1	i-2	ii-1	ii-2	p-1 /i-1	ii-2	ii-2	ii-2
Overcoating	I-1	I-1	I-1	I-1	I-1	I-2	I-3	I-4
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	3H	3H	3H	3H	3H	3H	3H	3H
Resistance to alkalis	good	good	good	good	good	good	good	good
Resistance to chemicals	good	good	good	good	good	good	good	good
Resistance to moisture	good	good	good	good	good	good	good	good
Resistance to climate	good	good	good	good	good	good	good	good
Resistance to water	good	good	good	good	good	good	good	good
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 15

Examples	9	10	11	12	13	14	15	16
Undercoating	ii-2							
Overcoating	I-5	I-6	I-7	I-8	I-9	II-1	II-2	II-3
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	3-	3-	3-	3-	3H	2H	2H	2H
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 16

Examples	17	18	19	20	21	22	23	24
Undercoating	ii-2	ii-2	ii-2	ii-2	ii-2	ii-2	ii-3	ii-2
Overcoating	II-4	II-5	II-6	II-7	II-8	II-9	II-10	II-11
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	2H	2H	2H	2H	2H	2H	3H	3H
Resistance to alkalis	good	good						

EP 1 022 318 A2

Table 16 (continued)

Examples	17	18	19	20	21	22	23	24
Undercoating	ii-2	ii-2	ii-2	ii-2	ii-2	ii-2	ii-3	ii-2
Overcoating	II-4	II-5	II-6	II-7	II-8	II-9	II-10	II-11
Resistance to chemicals	good	good						
Resistance to moisture	good	good						
Resistance to climate	good	good						
Resistance to water	good	good						
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 17

Examples	25	26	27	28	29	30	31	32
Undercoating	ii-2							
Overcoating	II-12	II-13	II-14	II-15	II-16	II-17	II-18	II-19
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	3H							
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

40 Examples 33 to 100

[0238] Each of the undercoating compositions was applied to the surface of a 50 µm-thick PET film to have an undercoating of 1 µm in thickness when dried. Then, a corresponding type of the overcoating composition was applied to the undercoating to have an overcoating of 0.1 µm in thickness when dried. After cured, a resultant coating film was analyzed for a variety of the characteristics.

[0239] The result of the analysis is shown in Tables 18 to 26.

Table 18

Examples	33	34	35	36	37	38	39	40
Undercoating	ii-4							
Overcoating	I-10	I-11	I-12	I-13	I-14	I-15	I-16	I-17
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	4H							
Resistance to alkalis	good							

EP 1 022 318 A2

Table 18 (continued)

Examples	33	34	35	36	37	38	39	40
Undercoating	ii-4							
Overcoating	I-10	I-11	I-12	I-13	I-14	I-15	I-16	I-17
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 19

Examples	41	42	43	44	45	46	47	48
Undercoating	ii-4	ii-4	ii-4	ii-4	ii-5	ii-5	ii-5	ii-5
Overcoating	I-18	I-19	I-20	I-21	I-10	I-12	I-14	I-16
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	4H	4H	4H	4H	4H	3H	4H	3H
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 20

Examples	49	50	51	52	53	54	55	56
Undercoating	ii-5	ii-5	ii-5	ii-6	ii-6	ii-6	ii-6	ii-6
Overcoating	I-17	I-19	I-21	I-10	I-11	I-12	I-13	I-14
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	4H							
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							

EP 1 022 318 A2

Table 20 (continued)

Examples	49	50	51	52	53	54	55	56
Undercoating	ii-5	ii-5	ii-5	ii-6	ii-6	ii-6	ii-6	ii-6
Overcoating	I-17	I-19	I-21	I-10	I-11	I-12	I-13	I-14
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

10

Table 21

Examples	57	58	59	60	61	62	63	64
Undercoating	ii-6	ii-6	ii-6	ii-6	ii-7	ii-7	ii-7	ii-7
Overcoating	I-15	I-16	I-17	I-18	I-10	I-11	I-13	I-15
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	4H							
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

15

20

25

30

35

Table 22

Examples	65	66	67	68	69	70	71	72
Undercoating	ii-7	ii-7	ii-7	ii-7	ii-8	ii-8	ii-8	ii-8
Overcoating	I-16	I-17	I-18	I-20	I-10	I-15	I-18	I-21
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	4H							
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

40

45

50

55

Table 23

Examples	73	74	75	76	77	78	79	80
Undercoating	ii-9							
Overcoating	I-10	I-13	I-14	I-15	I-17	I-18	I-19	I-21
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	4H							
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 24

Examples	81	82	83	84	85	86	87	88
Undercoating	ii-4	ii-5						
Overcoating	II-19	II-20	II-21	II-22	II-23	II-24	II-25	II-19
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	3-	3-	3-	3-	3H	2H	3H	3H
Resistance to alkalis	good							
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

Table 25

Examples	89	90	92	92	93	94	95	96
Undercoating	ii-5	ii-6	ii-6	ii-6	ii-6	ii-6	ii-7	ii-7
Overcoating	II-24	II-19	II-20	II-21	II-24	II-25	II-19	II-24
Adhesion (piece/100)	100	100	100	100	100	100	100	100
Pencil hardness	2H	3H	3H	3H	3H	2H	3H	3H
Resistance to alkalis	good							

EP 1 022 318 A2

Table 25 (continued)

Examples	89	90	92	92	93	94	95	96
Undercoating	ii-5	ii-6	ii-6	ii-6	ii-6	ii-6	ii-7	ii-7
Overcoating	II-24	II-19	II-20	II-21	II-24	II-25	II-19	II-24
Resistance to chemicals	good							
Resistance to moisture	good							
Resistance to climate	good							
Resistance to water	good							
Resistance to fouling	◎	◎	◎	◎	◎	◎	◎	◎
Transparency	○	○	○	○	○	○	○	○

15

Table 26

Examples	97	98	99	100
Undercoating	ii-7	ii-8	ii-9	ii-9
Overcoating	II-25	II-19	II-19	II-24
Adhesion (piece/100)	100	100	100	100
Pencil hardness	2H	3H	3H	3H
Resistance to alkalis	good	good	good	good
Resistance to chemicals	good	good	good	good
Resistance to moisture	good	good	good	good
Resistance to climate	good	good	good	good
Resistance to water	good	good	good	good
Resistance to fouling	◎	◎	◎	◎
Transparency	○	○	○	○

20

25

30

35

40 Examples 101 to 120

[0240] Each of the undercoating compositions was applied to the corresponding substrate listed in Table 27 and dried. Then, a corresponding type of the overcoating composition was applied to the undercoating and dried for curing. A resultant coating film was analyzed for the adhesion and the resistance to climate. The result of the analysis is shown in Table 27.

50

55

Table 27

	Substrate	Undercoat (thickness in μm)	Overcoat (thick- ness in μm)	Adhesion	Resistance to climate	Resistance to fouling
Examples						
101	Acryl film (50μm thick)	ii-2(1)	I-1(0.2)	100	good	○
102		ii-2(1)	II-1(0.2)	100	good	○
103		ii-4(1)	I-17(0.1)	100	good	○
104		ii-5(1)	I-17(0.1)	100	good	○
105		ii-9(1)	I-17(0.1)	100	good	○
106		ii-9(1)	I-18(0.1)	100	good	○
107		ii-4(1)	II-19(0.1)	100	good	○
108		ii-5(1)	II-24(0.1)	100	good	○
109		ii-7(1)	II-25(0.1)	100	good	○
110		ii-9(1)	II-25(0.1)	100	good	○
111	PC film	ii-2(0.5)	I-1(0.2)	100	good	○
112		ii-2(0.5)	II-1(0.2)	100	good	○
113		ii-4(1)	I-10(0.1)	100	good	○
114		ii-6(1)	I-16(0.1)	100	good	○
115		ii-7(1)	I-17(0.1)	100	good	○
116		ii-9(1)	I-18(0.1)	100	good	○
117		ii-4(1)	II-19(0.1)	100	good	○
118		ii-6(1)	II-25(0.1)	100	good	○
119		ii-7(1)	II-20(0.1)	100	good	○
120		ii-9(1)	II-21(0.1)	100	good	○

Effect of the Invention

[0241] The composition according to the present invention enables to provide a film which is improved in the resistance to fouling, the durability, and the transparency and has a photocatalytic function. The photocatalyst coating film made of the composition of the present invention can successfully be used preferably as a window film or more particularly as an automobile window film, a structure or house window film, a protective film, an UV barrier film, and an anti-fogging film.

[0242] A film which is improved in the resistance to fouling, the durability, and the transparency and has a photocatalytic function is provided by a method of making photocatalyst contained coating layers which comprises the steps of forming on the surface of an organic film an (under) coating of a coating composition composed mainly of at least one component (a) selected from organosilanes, hydrolyzates of the organosilanes, and condensates of the organosilanes and, when desired, a polymer component (b) having a silyl group containing silicon atoms bonded with a hydrolytic group and/or a hydroxy group, and forming on the (under)coating an (over)coating of a coating composition composed mainly of the component (a), a photocatalyst and, if desired, the component (b).

Claims

1. A method of making photocatalyst contained coating layers comprising the steps of providing a coating layer of the composition (i) or (ii) on the surface of an organic film and providing a coating layer of the composition (I) or (II) on the coating layer, wherein:

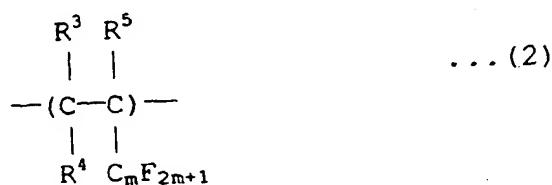
5 the composition (i) is a coating composition containing at least one component (a) selected from organosilanes, hydrolyzates of the organosilanes, and condensates of the organosilanes represented by Formula 1



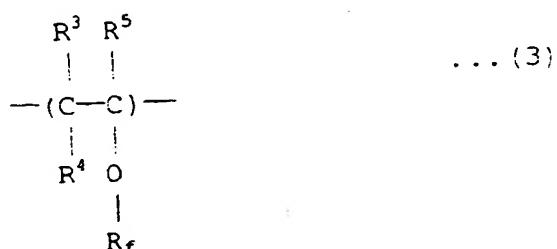
10 (wherein, R^1 is a monovalent organic group having 1 to 8 carbon atoms: when two exist, they are either identical to or different from each other; R^2 is an alkyl group having 1 to 5 carbon atoms or an acyl group having 1 to 6 carbon atoms: when two exist, they are either identical to or different from each other; and n is an integer ranging from 0 to 2);

15 the composition (ii) is a coating composition containing the component (a) and a polymer component (b) having a silyl group having silicon atoms bonded with a hydrolytic group and/or a hydroxy group; and the composition (I) is a coating composition containing the component (a) and a photocatalyst (c); and the composition (II) is a coating composition containing the component (a), the component (b), and the photocatalyst (c).

20 2. A method of making photocatalyst contained coating layers according to claim 1, wherein the component (b) in the composition (ii) is at least a substance selected from polymers having a structural unit expressed by Formula 2



25 (wherein, R^3 to R^5 are C_mY_{2m+1} , m is an integer ranging from 0 to 5, and Y is selected from F, H, and Cl separately) and/or a structural unit expressed by Formula 3



30 (wherein, R_f is an alkyl group or an alkoxyalkyl group having fluorine atoms and R^3 to R^5 are analogous to those in Formula 2 or may be modified without departing from the term of analogy), and simultaneously having a silyl group having silicon atoms bonded with a hydrolytic group and/or a hydroxy group.

35 3. A method of making photocatalyst contained coating layers according to claim 1, wherein the component (b) in the composition (II) is at least a substance selected from polymers having a structural unit expressed by Formula 2 and/or a structural unit expressed by Formula 3 and simultaneously having a silyl group containing silicon atoms bonded with a hydrolytic group and/or a hydroxy group.

40 4. A method of making photocatalyst contained coating layers according to claim 1, wherein the composition (i) or (ii) further contains an ultraviolet light absorber (d).

45 5. A method of making photocatalyst contained coating layers according to claim 1, wherein the composition (I) or (II) further contains a colloidal silica and/or a colloidal alumina (e).

50 6. A method of making photocatalyst contained coating layers according to claim 1, wherein the composition (i) or (ii) further contains a colloidal silica and/or a colloidal alumina (e).

7. A method of making photocatalyst contained coating layers according to claim 1, wherein the component (a) in at least one of the compositions (i), (ii), (I), and (II) has partially or at its entirety an SiO bond and a siloxane oligomer of which the weight average molecular weight ranges from 300 to 100,000.

5 8. A method of making photocatalyst contained coating layers according to claim 1, wherein at least one of the compositions (i), (ii), (I), and (II) has a curing catalyst (f).

9. A method of making photocatalyst contained coating layers according to claim 7, wherein the curing catalyst (f) is an organic metal compound.

10 10. A method of making photocatalyst contained coating layers according to claim 1, wherein at least one of the compositions (i), (ii), (I), and (II) contains at least a compound selected from β -diketones and β -ketoesters (g) expressed by Formula 4



(wherein, R^6 is a monovalent hydrocarbon group having 1 to 6 carbon atoms, such as methyl, ethyl, n-propyl, i-propyl, n-butyl, sec-butyl, t-butyl, n-pentyl, n-hexyl, cyclohexyl, or phenyl, and R^7 is either a monovalent hydrocarbon group having 1 to 6 carbon atoms, like R^6 , or an alkoxy group having 1 to 16 carbon atoms, such as methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, sec-butoxy, t-butoxy, lauryloxy, or stearyloxy).

20 11. A photocatalyst coating film produced by providing coating layers on the surface of an organic film with the use of a method of making photocatalyst contained coating layers defined in any of claims 1 to 10.

25

30

35

40

45

50

55

THIS PAGE BLANK (USPTO)

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets

(11)



EP 1 022 318 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
28.11.2001 Bulletin 2001/48

(51) Int Cl.7: C09D 183/04, C08J 7/04,
C08K 3/00

(43) Date of publication A2:
26.07.2000 Bulletin 2000/30

(21) Application number: 00100911.7

(22) Date of filing: 18.01.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 19.01.1999 JP 1112199
21.05.1999 JP 14240999

(71) Applicant: JSR Corporation
Tokyo 104-8410 (JP)

(72) Inventors:
• Kanamori, Tarou, c/o JSR Corporation
Tokyo 104-8410 (JP)

• Shimada, Mibuko, c/o JSR Corporation
Tokyo 104-8410 (JP)
• Sakagami, Toshinori, c/o JSR Corporation
Tokyo 104-8410 (JP)

(74) Representative:
Pellmann, Hans-Bernd, Dipl.-Ing. et al
Patentanwaltsbüro
Tiedtke-Bühling-Kinne & Partner
Bavariaring 4-6
80336 München (DE)

(54) Method of making coating layers containing photocatalyst and a photocatalyst coating film formed thereby

(57) A film which is improved in the resistance to fouling, the durability, and the transparency and has a photocatalytic function is provided by a method of making photocatalyst contained coating layers which comprises the steps of forming on the surface of an organic film an (under)coating of a coating composition composed mainly of at least one component (a) selected

from organosilanes, hydrolyzates of the organosilanes, and condensates of the organosilanes and, when desired, a polymer component (b) having a silyl group-containing silicon atoms bonded with a hydrolytic group and/or a hydroxy group, and forming on the (under)coating an (over)coating of a coating composition composed mainly of the component (a), a photocatalyst and, if desired, the component (b).

European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 00 10 0911

DOCUMENTS CONSIDERED TO BE RELEVANT			CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	
P, X	EP 0 916 411 A (MATSUSHITA ELECTRIC WORKS LTD) 19 May 1999 (1999-05-19) * claims 1-26 * * page 8, line 31 - page 9, line 39 * & WO 98 25711 A 18 June 1998 (1998-06-18) ---	1	C09D183/04 C08J7/04 C08K3/00
X		1	
P, X	EP 0 923 988 A (NIPPON SODA CO) 23 June 1999 (1999-06-23) * claims 1-22 * * page 3, line 20 - line 55 * * page 4, line 31 - page 5, line 3 * * page 5, line 45 - page 7, line 15 * * page 8, line 15 - line 19 * * page 12, line 40 - page 14, line 1 * & WO 97 00134 A 3 January 1997 (1997-01-03) ---	1	
X		1	
E	EP 0 987 317 A (TOTO LTD) 22 March 2000 (2000-03-22) * claims 1,8-14,19-21 * * page 2, line 20 - line 48 * * page 5, line 25 - page 6, line 3 * * page 6, line 51 - page 8, line 37 * * page 10, line 10 - page 11, line 29 * & WO 98 55573 A 10 December 1998 (1998-12-10) ---	1	
E		1	
X		1	TECHNICAL FIELDS SEARCHED (Int.Cl.7) C08J C09D C08K
E	EP 1 029 659 A (NIPPON SODA CO) 23 August 2000 (2000-08-23) * claims 1-22 * * page 11; example 1 * & WO 99 24253 A 20 May 1999 (1999-05-20) ---	1	
P, X		1	
		-/-	
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search	Examiner	
THE HAGUE	3 October 2001	Depijper, R	
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 10 0911

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (IntCl.7)
A	EP 0 824 127 A (JAPAN SYNTHETIC RUBBER CO LTD) 18 February 1998 (1998-02-18) * claims 1-20 * * page 3, line 20 - line 25 * * page 3, line 42 - line 47 * * page 4, line 1 - line 25 * * page 9, line 40 - page 13, line 27 * * page 14, line 40 - line 55 * ----	1	
A	EP 0 869 156 A (TOTO LTD) 7 October 1998 (1998-10-07) * claims 1-29 * * page 16; example 7 *	1	
A	EP 0 633 064 A (FUJISHIMA AKIRA ; HASHIMOTO KAZUHITO (JP); ISHIHARA SANGYO KAISHA ()) 11 January 1995 (1995-01-11) * claims 1-12 * * page 3, line 26 - line 41 * * page 4, line 25 - line 32 *	1	
A	US 5 755 867 A (CHIKUNI MAKOTO ET AL) 26 May 1998 (1998-05-26) * claims 1-29 * * column 13 - column 14; example 1 *	1	TECHNICAL FIELDS SEARCHED (IntCl.7)
The present search report has been drawn up for all claims			
Place of search EPO FORM 1503/03 03/02 P04C01	Date of completion of the search 3 October 2001	Examiner Depijper, R	
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			
T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 00 10 0911

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
 The members are as contained in the European Patent Office EDP file on
 The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

03-10-2001

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0916411	A	19-05-1999	JP 3182107 B2 JP 10225658 A EP 0916411 A1 US 6165619 A WO 9825711 A1	03-07-2001 25-08-1998 19-05-1999 26-12-2000 18-06-1998
EP 0923988	A	23-06-1999	JP 9310039 A EP 0923988 A1 JP 3038599 B2 US 6228480 B1 WO 9700134 A1	02-12-1997 23-06-1999 08-05-2000 08-05-2001 03-01-1997
EP 0987317	A	22-03-2000	JP 10337526 A AU 7550598 A BR 9810241 A EP 0987317 A1 CN 1259162 T WO 9855573 A1 JP 1105006 A	22-12-1998 21-12-1998 05-09-2000 22-03-2000 05-07-2000 10-12-1998 23-02-1999
EP 1029659	A	23-08-2000	JP 11138686 A EP 1029659 A1 WO 9924253 A1 JP 11267516 A	25-05-1999 23-08-2000 20-05-1999 05-10-1999
EP 0824127	A	18-02-1998	JP 10060280 A EP 0324127 A2 US 5905109 A	03-03-1998 18-02-1998 18-05-1999
EP 0869156	A	07-10-1998	US 6090489 A AU 1170797 A EP 0869156 A1 JP 2917525 B2 AU 718733 B2 AU 5014096 A BR 9607868 A CA 2215925 A1 CA 2241059 A1 CN 1209153 A CN 1184498 A DE 19681289 T0 DE 29623901 U1 EP 0816466 A1 JP 10146251 A JP 9231821 A WO 9629375 A1	18-07-2000 17-07-1997 07-10-1998 12-07-1999 20-04-2000 08-10-1996 30-06-1998 26-09-1996 03-07-1997 24-02-1999 10-06-1998 16-04-1998 06-07-2000 07-01-1998 02-06-1998 05-09-1997 26-09-1996

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 00 10 0911

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

03-10-2001

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0869156 A		WO 9723572 A1	03-07-1997
		JP 9229585 A	05-09-1997
		JP 9231819 A	05-09-1997
		JP 9226040 A	02-09-1997
		JP 9225392 A	02-09-1997
		JP 9225396 A	02-09-1997
		JP 9225385 A	02-09-1997
		JP 9227178 A	02-09-1997
		JP 10140046 A	26-05-1998
		JP 3189682 B2	16-07-2001
		JP 9225388 A	02-09-1997
		JP 9228072 A	02-09-1997
		JP 9228073 A	02-09-1997
		JP 9225276 A	02-09-1997
		JP 9224490 A	02-09-1997
		JP 9225263 A	02-09-1997
		JP 9229493 A	05-09-1997
		JP 9231849 A	05-09-1997
		JP 2943763 B2	30-05-1999
		JP 10156999 A	16-06-1998
		JP 9225021 A	02-09-1997
		JP 9226695 A	02-09-1997
		JP 9230104 A	05-09-1997
		JP 9232096 A	05-09-1997
		JP 3129194 B2	29-01-2001
		JP 9226054 A	02-09-1997
		JP 9228307 A	02-09-1997
		JP 9225389 A	02-09-1997
		JP 11070613 A	16-03-1999
		JP 111537 A	
EP 0633064 A	11-01-1995	AT 174812 T	15-01-1999
		AU 676299 B2	06-03-1997
		AU 6481094 A	05-01-1995
		CA 2126418 A1	29-12-1994
		CN 1101591 A , B	19-04-1995
		CN 1256172 A	14-06-2000
		DE 69415420 D1	04-02-1999
		DE 69415420 T2	02-06-1999
		DK 633064 T3	23-08-1999
		EP 0633064 A1	11-01-1995
		EP 0875289 A1	04-11-1998
		JP 2918787 B2	12-07-1999
		JP 7171408 A	11-07-1995
		US 5547823 A	20-08-1996
		US 6277346 B1	21-08-2001

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 00 10 0911

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

03-10-2001

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5755867 A	26-05-1998 JP	9227829 A	02-09-1997

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- BLACK BORDERS**
- IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- FADED TEXT OR DRAWING**
- BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- SKEWED/SLANTED IMAGES**
- COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- GRAY SCALE DOCUMENTS**
- LINES OR MARKS ON ORIGINAL DOCUMENT**
- REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.

THIS PAGE BLANK (USPTO)